

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 22, 2005, 10:02:08 ; Search time 178 Seconds

(without alignment)
1668.573 Million cell updates/sec

Title: US-10-629-951-2

Perfect score: 3055
Sequence: 1 MGTGLSLSLGDRGAPTV.....HKLNKTHDMLNENKLSLS 580

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1612378 seqs, 512079187 residues

Total number of hits satisfying chosen parameters: 1612378

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: uniprot_03.*
2: uniprot_trembl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	3055	100.0	580	1	MBD4_HUMAN
2	1821.5	59.6	554	1	MBD4_MOUSE
3	880.5	28.8	416	2	Q919F1
4	344	11.3	419	2	Q9SFC1
5	338	11.1	407	2	Q84WT3
6	238.5	7.8	524	2	Q71T77
7	220.5	7.2	467	2	Q9YGC6
8	213	7.0	682	2	Q7SCQ2
9	208.5	6.8	486	1	MEC2_HUMAN
10	208.5	6.8	498	2	Q6OH99
11	208.5	6.8	516	2	Q7Z384
12	207.5	6.6	486	1	MEC2_MACFA
13	202.5	6.6	344	2	Q42403
14	201	6.6	492	1	MEC2_RAT
15	198.5	6.5	484	1	MEC2_MOUSE
16	156	5.1	2473	2	Q7RN41
17	149.5	4.9	2212	2	Q81T6
18	148.5	4.9	352	2	Q8L876
19	148.5	4.9	1679	2	Q7RCP2
20	146	4.8	702	2	Q9SFP2
21	146	4.8	1643	2	Q8HZQ1
22	145	4.7	267	2	Q8AYT1
23	145	4.7	1301	1	SAC3_YEAST
24	145	4.7	3418	1	BRC2_HUMAN
25	144	4.7	283	2	Q8AYP2
26	143.5	4.7	782	2	Q25875
27	141.5	4.6	782	2	Q25730
28	141.5	4.6	782	2	Q26007
29	140.5	4.6	782	2	Q9U414
30	140.5	4.6	782	2	Q9U430
31	140.5	4.6	881	2	Q7RK82

32	140.5	4.6	1030	2	Q924D9	Q924D9 meriones sh
33	138.5	4.5	500	1	GAR2_SCHPO	P41891 schizosacch
34	138.5	4.5	782	2	Q9U479	Q9U479 plasmodium
35	138.5	4.5	1047	2	Q7REU9	Q7REU9 plasmodium
36	137.5	4.5	326	2	Q869D5	Q869D5 branchiosteo
37	137.5	4.5	412	2	Q6MFI1	Q6MFI1 neureospora
38	137.5	4.5	782	2	Q26104	Q26104 plasmodium
39	137.5	4.5	2209	2	Q9U0G6	Q9U0G6 plasmodium
40	137	4.5	400	2	Q8LK06	Q8LK06 zea mays (m
41	137	4.5	560	1	SET4_YEAST	P42948 saccharomyc
42	137	4.5	707	2	Q7RKH5	Q7RKH5 plasmodium
43	137	4.5	1713	2	Q7T550	Q7T550 mus musculu
44	137	4.5	1813	2	Q75X83	Q75X83 helicobacte
45	136.5	4.5	598	1	CYL1_HUMAN	P35663 homo sapien

ALIGNMENTS

RESULT 1
MBD4_HUMAN STANDARD; PRT; 580 AA.
AC Q95243; Q7Z4T3; Q96F09;
DT 25-OCT-2004 (Rel. 45, Created)
DT 25-OCT-2004 (Rel. 45, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Methyl-Cpg binding protein 4 (BC 3.2.2.-) (Methyl-Cpg binding domain
DE protein 4) (Methyl-Cpg binding endonuclease 1) (Mismatch-specific DNA
DE N-glycosylase).
GN Name=MBD4; Synonyms=MBD1;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP MEDLINE=96449942; PubMed=9774669;
RA Hendrich B., Bird A.;
RT "Identification and characterization of a family of mammalian methyl-
RT Cpg binding proteins.";
RL Mol. Cell. Biol. 18:6538-6547(1998).
RN [2]
RP SEQUENCE FROM N.A. (ISOFORM 1).
RX MEDLINE=99373255; PubMed=10441743;
RA Hendrich B., Abbott C., McQueen H., Chambers D., Cross S.H., Bird A.;
RT "Genomic structure and chromosomal mapping of the murine and human
RT mbd1, mbd2, mbd3, and mbd4 genes.";
RL Mamm. Genome 10:906-912(1999).
RN [3]
RP SEQUENCE FROM N.A. (ISOFORM 1), FUNCTION, AND INTERACTION WITH MLH1.
RC TISSUE=Petal brain;
RX MEDLINE=99199294; PubMed=10097147; DOI=10.1073/pnas.96.7.3969;
RA Bellacosa A., Cicchilitti L., Schepis F., Riccio A., Yeung A.T.,
RA Matsumoto Y., Golemis E.A., Genuardi M., Neri G.;
RT "MBD1, a novel human methyl-Cpg-binding endonuclease, interacts with
RT DNA mismatch repair protein MLH1.";
RL Proc. Natl. Acad. Sci. U.S.A. 96:3969-3974(1999).
RN [4]
RP SEQUENCE FROM N.A. (ISOFORM 3).
RC TISSUE=Lung;
RA Guo J.H., Chen L., Yu L.;
RT Submitted (Jul-2002) to the EMBL/Genbank/DBJ databases.
RN [5]
RP SEQUENCE FROM N.A. (ISOFORM 1), AND VARIANTS SER-273; PRO-342; LYS-346
RP AND HIS-568.
RA Rieder M.J., Braun A.C., Montoya M.A., Chung M.-W., Nguyen C.P.,
RA Nguyen D.A., Livingston R.J., Poel C.L., Robertson P.D.,
RA Schackwitz W.S., Sherwood J.K., Wiltz L.A., Nickerson D.A.;
RT "NIHES-SNP, environmental genome project, NIHES ES15478, Department
RT of Genome Sciences, Seattle, WA (URL: <http://egp.gs.washington.edu>).";
RL Submitted (MAR-2002) to the EMBL/Genbank/DBJ databases.
RN [6]
RP SEQUENCE FROM N.A. (ISOFORM 2).

Query Match	Similarity	Score	DB 1	Length	580;
Best Local	Similarity	100.0%;	Pred. No. 5.3e-178;		
Matches	580;	Conservative	0;	Mismatches	0;
				Indels	0;
				Gaps	0;
EMBL; AF120999; AAD50374.1; -					
DR EMBL; AF120997; AAD50374.1; JOINED.					
DR EMBL; AF120998; AAD50374.1; JOINED.					
DR EMBL; AF114784; AAD22195.1; -					
DR EMBL; AF532602; AAP97338.1; -					
DR EMBL; AF494057; AAM00008.1; -					
DR EMBL; CR450305; CAG29301.1; -					
DR EMBL; BC011752; AAH11752.1; -					
DR HSSP; Q922D7; INGN.					
DR IntAc; Q95243; -					
DR Genew; HGNC:6919; MBD4.					
DR H-InVD; HIX0003669; -					
DR Reactome; O95243; -					
DR MIM; 603574; -					
DR GO; GO:0005634; C:nucleus; TNS.					
DR GO; GO:0004520; F:endodeoxyribonuclease activity; TNS.					
DR GO; GO:0003696; F:stellite DNA binding; TNS.					
DR GO; GO:0006281; P:DNA repair; TNS.					
DR InterPro; IPR003265; Endo_3c.					
DR InterPro; IPR001739; Methyl-CpG_bind.					
DR Pfam; PF00730; Hhh-GPD, 1.					
DR Pfam; PF01429; MBD, 1.					
DR SMART; SM00391; MBD, 1.					
DR PROSITE; PS50982; MBD, 1.					
KM Alternative splicing; DNA repair; DNA-binding; Hydrolyase;					
KW Nuclear protein; Polymorphism.					
FT DOMAIN 76 148					
FT Hhh-GPD.					
FT By similarity.					
FT Missing (in isoform 2).					
FT /FtId=VSP_010816.					
FT Ky -> Ap (in isoform 3).					
FT /FtId=VSP_010817.					
FT Missing (in isoform 3).					
FT /FtId=VSP_010818.					
FT A -> S (in dbSNP:10342).					
FT /FtId=VAR_019357.					
FT A -> T (in dbSNP:10342).					
FT /FtId=VAR_019514.					
FT S -> P (in dbSNP:2307289).					
FT /FtId=VAR_019515.					
FT E -> K (in dbSNP:146693).					
FT /FtId=VAR_019359.					
FT I -> T (in dbSNP:2307298).					
FT /FtId=VAR_019515.					
FT D -> H (in dbSNP:2307293).					
FT /FtId=VAR_019360.					
FT B16FB21A34B85F CRC64;					
SEQUENCE 580 AA; 66050 MW; B16FB21A34B85F CRC64;					

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QY 301 SVTSENSLVKKKERSLSGSGNFCSEOKTSGIINKFCSAKSEHNEKEDPFLSEELGT 360
DB 301 SVTSENSLVKKKERSLSGSGNFCSEOKTSGIINKFCSAKSEHNEKEDPFLSEELGT 360
QY 361 KVEVERKEHLHTDILKXGSEMDNNSCPTRKDPFTEKIFQEDPTIPTQIERRKTSLYPSS 420
DB 361 KVEVERKEHLHTDILKXGSEMDNNSCPTRKDPFTEKIFQEDPTIPTQIERRKTSLYPSS 420
QY 421 KYNKALSPRRKAKFKWTPPRSPNLVOETLPHDPWKLILATIFLANTSGMAIPVLWK 480
DB 421 KYNKALSPRRKAKFKWTPPRSPNLVOETLPHDPWKLILATIFLANTSGMAIPVLWK 480
QY 481 FLEKPPSAEVARTAWRDVSELKVLGLYDLAKTIKVSDEDTLTKOMKYPTEILHGIGKY 540
DB 481 FLEKPPSAEVARTAWRDVSELKVLGLYDLAKTIKVSDEDTLTKOMKYPTEILHGIGKY 540
QY 541 GNDSTRIFCVNEMKOVHPEDHLKXKHPDLWENHEKLSLS 580
DB 541 GNDSTRIFCVNEMKOVHPEDHLKXKHPDLWENHEKLSLS 580

RESULT 2
MED4_MOUSE STRAND: PRT; 554 AA.
ID MED4_MOUSE Q922D7; Q792D2; Q8R3R3;
AC 25-OCT-2004 (Rel. 45, Created)
DT 25-OCT-2004 (Rel. 45, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Methyl-Cpg binding protein 4 (BC 3.2.2.-) (Methyl-Cpg binding domain
  protein 4) (Mismatch-specific DNA N-glycosylase).
GN Name=Mbd4;
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A., FUNCTION, AND SUBCELLULAR LOCATION.
RX MEDLINE=98449942; PubMed=9774669;
RA Hendrich B., Bird A.;
RT "Identification and characterization of a family of mammalian methyl-
  Cpg binding proteins."
RT Mol. Cell. Biol. 18:6538-6547 (1998).
RN [2]
RP SEQUENCE FROM N.A.
RX STRAIN=129;
RC MEDLINE=99373255; PubMed=10441743;
RA Klausner R.D., Collins F.S., Wagner L., Shennan C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buettow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Heist L.,
RA Ditscheko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Donald M.F., Casavant T.L., Schaefer T.E.,
RA Brownstein M.J., Ueda T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.D., Abramson R.D., Mullaly S.J.,
RA Bosak S.A., McMan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny K.C., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahy J., Heiton E., Kettelman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Buterfield Y.S.N., Krzywinski M.I., Skalska U., Smalins D.E.,
RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
  and mouse cDNA sequences."
RT Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).

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RN [4]
RP X-RAY CRYSTALLOGRAPHY (2.1 ANGSTROMS) OF 411-554.
RX PubMed12456671; DOI=10.1074/jbc.M210848200;
RA Wu P., Qiu C., Shail A., Zhang X., Bhagwat A.S., Cheng X.,
RT "Mismatch repair in methylated DNA. Structure and activity of the
  mismatch-specific thymine glycosylase domain of methyl-Cpg-binding
  protein MBD4."
RT J. Biol. Chem. 278:5285-5291 (2003).
RL J. Biol. Chem. 278:5285-5291 (2003).
CC -1- FUNCTION: Mismatch-specific DNA N-glycosylase involved in DNA
  repair. Has thymine glycosylase activity and is specific for G:T
  mismatches within methylated and unmethylated Cpg sites. Can also
  remove uracil or 5-fluorouracil in G:U mismatches. Has no lyase
  activity. Was first identified as methyl-Cpg-binding protein.
CC -1- SUBUNIT: Interacts with MTH1 (By similarity).
CC -1- SUBCELLULAR LOCATION: Nuclear. In discrete foci.
CC -1- SIMILARITY: Contains 1 methyl-Cpg-binding (MBD) domain.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
  between the Swiss Institute of Bioinformatics and the EMBL outstation -
  the European Bioinformatics Institute. There are no restrictions on its
  use by non-profit institutions as long as its content is in no way
  modified and this statement is not removed. Usage by, and for commercial
  entities requires a license agreement (see http://www.isb-sib.ch/announce/
  or send an email to license@isb-sib.ch).
CC -----
DR EMBL AF072249; AAC68878.1; -
DR EMBL AF120996; AAD56595.1; -
DR EMBL BC024812; AAH24812.1; -
DR PDB: 1NGN; X-ray; A=400-554.
DR MGD: MGI:1333850; Mbd4.
DR InterPro: IPR001739; Endo_3c.
DR InterPro: IPR001739; Methyl-Cpg_bind.
DR Pfam: PF00730; HNH-GPD; 1.
DR Pfam: PF01429; MBD; 1.
DR SMART: SM00391; MBD; 1.
DR PROSITE: PS50982; MBD; 1.
KW 3D-structure; DNA repair; DNA-binding; Hydroxylase; Nuclear protein.
FT DOMAIN 63 135 MBD.
FT DOMAIN 435 498 HNH-GPD.
FT ACT_SITE 534 534 By similarity.
FT CONFLICT 129 129 N -> D (in Ref. 3).
SQ SEQUENCE 554 AA; 62577 MW; 792D37CB180291F5 CRC64;

Query Match 59.6%; Score 1821.5; DB 1; Length 554;
Best Local Similarity 66.2%; Pred. No. 7.6e-103;
Matches 384; Conservative 49; Mismatches 116; Indels 31; Gaps 11;

QY 6 LESLSIGD---RGAPVTYSSERLVPDPNDLRKEDVAMELERVGEDEBQMMIKRSSSECN 62
DB 1 MESPNLGDNRVVG-----ESLVPDPDPMDRCKEDIVAGLGVGEDGKDLVI--SSERS 50
QY 63 PLLQEPPIASQPGATAGTECRKSVPCGWERVVKQLFGKTAGRPDPVYFISPGILKFRSKS 122
DB 51 SLQLEPTAST-LSSTTATGEHKKVPFCGWERVVKQLSGKTAKFPVYFISPGILKFRSKR 109
QY 123 SLANTLHNGEISLPEDPDPFVVLKRGKIKSKYKDCSMAALTSIH.QONSONNSNMULRPS 182
DB 110 SLANTYLLKNGEFTLPDEPNFVLPKGSINPGYKQSLAALTSLOPNETDVSQKULKTR 169
QY 183 KKKQVFPMPSSSSSELSQSGLSNFTSTHLLKKEDEGVDDVAFRKVRKPKGVYLLKGP 242
DB 170 KKKTDVLPSPSTSSPSSSGISNSNSACILIREHRDIDVDSEKRRKSKRVYLVKGR 229
QY 243 IKTKKXGCRKSCSGFVQSDSKRESVCKNDAESEPVAQSKQLDRVTCISDAGACBETSLV 302
DB 230 SKTKQKCKKSLBSSTQRRKRSVYQKVGADRELVPQSSQLNRTLCPADACA-RETVGL 288
QY 303 TEENSVLKKGRSLSSGNSFCSEOKTSGIINKFCSAKSEHNEKEDPFLSEELGTAV 362
DB 289 AGE-----EKSSPPGIDLCFIQVTSGTINKKFTSTEAAGANR-EQTFLESEELRKS- 338
QY 363 EVVERK--EHLHTDILKXGSEMDNNSCPTRKDPFTEKIFQEDPTIPTQIERRKTSLYPSS 420

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Db      339  --GDRKGAAHLHTGVLDQDSEMP--SSGQAKKHFTSE--TIQEDSIPRTQVEKRTSLYFSS 394

QY      421  KKNKEALSPRRKAKFKKMTPPRSPFNLYQETLFHDEWKLIIATIFLNRTSGKVALPYLWK 480
         : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db      395  KKNKEALSPRRKRSFKKMTPPRSPFNLYQETLFHDEWKLIIATIFLNRTSGKVALPYLWE 454

QY      481  FLEKTPSAEVAFTADWRDVSLLKPLGLYDLRAKTIYKSSDEYLTQQMKPIELHAGIGKY 540
         : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db      455  FLEKTPSAEVAADWRDVSLLKPLGLYDLRAKTIYKSSDEYLTQQMKPIELHAGIGKY 514

QY      541  GNDSTYRIFCVNEMKQVHPEDHLKNTKCHDWLMENHEGLSTLS 580

Db      515  GNDSTYRIFCVNEMKQVHPEDHLKNTKCHDWLMENHEGLSTLS 554

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RESULT 3

ID	Q919F1	PRELIMINARY;	FRT;	416 AA.
AC	Q919F1.			
DT	01-OCT-2000 (TREMBLrel. 15, Created)			
DT	01-OCT-2000 (TREMBLrel. 15, Last sequence update)			
DT	01-OCT-2003 (TREMBLrel. 25, Last annotation update)			
DE	5-methylcytosine G/T mismatch-specific DNA glycosylase.			
OS	Gallus gallus (chicken).			
OC	Eukaryota; Metazoa; Chordata; Craniota; Vertebrata; Euteleostomi;			
OC	Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;			
OC	Gallus.			
OX	NCBI_TaxID=9031;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
FX	MEDLINE=20512562; PubMed=11058112; DOI=10.1093/nar/28.21.4157.			
RA	Zhu B., Zheng Y., Anglikar H., Schwarz S., Thiry S., Stegmann M.,			
RA	Joest U.-P.,			
RT	"5-Methylcytosine DNA glycosylase activity is also present in the			
RT	human MBD4 (G/T mismatch glycosylase) and in a related avian			
RT	sequence."			
RL	Nucleic Acids Res. 28:4157-4165(2000).			
DR	EMBL, AF257107; AAF68981.1; -.			
DR	HSSP, Q922D7; INGN.			
DR	InterPro, IPR011257; DNA_glycylase.			
SO	SEQUENCE 416 AA; 45454 MW; AAF70C6FF2133F2A CRC64;			

Query

Query Match	28.8%	Score 880.5	DB 2	Length 416
Best Local Similarity	78.2%	Pred. NO. 1.2e-45		
Matches 161; Conservative	20	Mismatches 21	Indels 4	Gaps 2

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OY 378 GGESEMDNNNS--PTKRDFFGEKIFQ--EDTPTQIERTKTSYLSFSKYNKALSPRRK 433
Db 210 RDSAADGDVSWPDSKKSFTAYVAPRGTSEAPPTQVDRKRTSPFSKSKSKEALSPRRK 269
OY 434 AFKKATPPRSPFNLYVOETLFHDPMKLLIATIPLNRTSGKMAIVLWKFLFKYPSAEVART 493
   ||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 270 AFRKRTTPRSPFNLYVOETLFHDPMKLLIATIFLNKTSKGMAIVLWELFKYSPSEVART 329
OY 494 ADMRDVSELKPLGLYIDURAKTIKFSDEYLTQWKYPIELHIGIGKXGDSYRIFCVNEW 553
   ||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 330 ADMKEMSELRLPLGLYALPRAKTIKFSDEYLNQWKYPIELHIGIGKXGDSYRIFCVNEW 369
OY 554 KQYHPEDKLNKYHDMWLNHEKLSL 579
Db 390 KEVQPODHRQLNLYHAWLNHEKLSV 415

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RESULT 4

ID	Q9SEF1	PRELIMINARY;	PRT;	419 AA.
AC	Q9SEF1;			
DT	01-MAY-2000	(TRENBLrel. 13, Created)		
DT	01-MAY-2000	(TRENBLrel. 13, Last sequence update)		
DT	01-JUN-2003	(TRENBLrel. 24, Last annotation update)		
DE	F17A17.27 protein.			
GN	Name=F17A17.27;			
OS	Arabidopsis thaliana (Mouse-ear cress).			

OC Eukaryot; Vitellidiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
 OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 OK NCBI_TaxID=3702;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Lin X., Kaul S., Town C.D., Benito M.-I., Creasy T.H., Haas B.,
 RA Rowning C.M., Koo H., Fujii C.Y., Ultebrack T.R., Barnstead M.E.,
 RA Bowman C.L., White O., Niernan W.C., Fraser C.M.;
 RL Submitted (Jan-2001) to the EMBL/GenBank/DBD databases.
 DR EMBL; AC013483; AAF21203.1; -.
 DR HSSP; Q9Z2D7; INGN.
 DR GO; GO:0006284; P:base-excision repair; IEA.
 DR InterPro; IPR011257; DNA_glycylase.
 DR InterPro; IPR003265; Endo_3c.
 DR Pfam; PF00730; HNH-GPD; 1-
 SQ SEQUENCE 419 AA; 47957 MW; BAF0BCA5A710C95 CRC64;

QY

QY	205	SNFTSTHLKKEDEGVDDVNRKVRKPKGKTYILKGIPI	-----	KTKTKGCRKSCGF	257
Db	88	SNLVSPLQADDDDSVDSHIERQCESEFHVVRKRVSPYQGS	-----	QSQSKKGC	140

[illegible]

RESULT 5

TO			
ID	084WT3	PRELIMINARY;	PRT, 407 AA.
AC	084WT3;		
DT	01-JUN-2003	(TREMBLrel. 24, Created)	
DT	01-JUN-2003	(TREMBLrel. 24, Last sequence update)	
DT	01-OCT-2003	(TREMBLrel. 25, Last annotation update)	
DE	Hypothetical protein At3g07930.		
GN	Name=At3g07930;		
OS	Arabidopsis thaliana (Mouse-ear cress).		
OC	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;		
OC	Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;		
OC	eurosids II; Brassicales; Brassicaceae; Arabidopsis.		
OX	NCBI_TaxID=3702;		
RN	[1]		
RP	SEQUENCE FROM N.A.		
RA	Yamada K., Chan M.M., Chang C.H., Dale J.M., Hsuan V.W., Lee J.M.,		
RA	Onodera C.S., Quach H.L., Tang C., Tottum M., Wong C., Wu H.C.,		
RA	Yu G., Yuan S., Carminci P., Chen H., Cheuk R., Hayashizaki Y.,		
RA	Ishida J., Jones T., Kamiya A., Kawai J., Kim C.J., Natsuoka M.,		
RA	Nguyen M., Palm C.J., Sakurai T., Satou M., Seki M., Shim P.,		
RA	Southwick A., Tzipi M.G., Wu T., Shinzaki K., Davis R.W., Ecker J.R.,		

RA Theologis A.;
 RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
 DR EMBL: BT002799; AA02623.1; -.
 DR HSSP: Q922D7; INGN.
 DR GO: GO:0006284; P:base-excision repair; IEA.
 DR InterPro: IPR011257; DNA_glycosylase.
 DR InterPro: IPR003465; Endo_3c.
 DR Pfam: PF00730; HNH-GPD; 1.
 DR Hypothetical protein.
 KM
 SQ SEQUENCE 407 AA; 46941 MW; DD758CD862EF54F CRC64;

Query Match 11.1%; Score 338; DB 2; Length 407;
 Best Local Similarity 26.6%; Pred. No. 1,4e-12;
 Matches 102; Conservative 51; Mismatches 155; Indels 76; Gaps 11;

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QY 251 RKSCGFWQ-SDSKESVYCNKADASEPVAQKSQLDRFVCI SDAGACGTTLSVTSENSL 309
DB 30 RRPDDDFIEVDENSNFALFKEDDEK-----NRDLGLVDDGSTNLVLCQHDGSL 80
QY 310 YKKKERSLS---SGSNFCSEQR-----TSGIINKFCSAKDEHNEKTEDTFLBSE 356
DB 81 EKDNNSLDDLFSGFYKGVRRKRKDDFGSITTSNLSVPOIADD--DDSVDSHIERO 137
QY 357 EIGTKVEYVERK-----EHLHTDLKRGSEMDN-NCSPTRKDTGTEKI---FOEDT 403
DB 138 EESKVOAKVPRVSPYFOASTISQCDSDIVSSQSGNRKSGSSKQVKARVSPYQEST 197
QY 404 IP-----RTQIERKRTSLYF-----SSKYNKEALSP----- 430
DB 198 VEEQNPQAKGLRNTFKVVKVSRFYFADGIVQNESQKESRVRKTPYSPVLSQKTD 257
QY 431 ---RRKAFKKWTTPRSFNNLVQETLFHDPMKLLATIFLARTSGKMLPVLWKELEKP 486
DB 258 DYLRKTPNTWVPPSPCNLLQEDHMDPMRVLVTCMLNTKSGAQTRGVISDLFGICT 317
QY 487 SAEVARTADWRDVSSELKPLGLYDLRAKTIYKPSDYILTKQKPIYELHGLICRYGNDYR 546
DB 318 DAKTATEVEVEEIEENIKPLQKRTKMIORLSLEYLOESWTHVQLHGKTAADAYA 377
QY 547 IFCVNEWKQVHPEDHLKYNKHWL 570
DB 378 IFGNGMWRKVRPDHMLNTYMDL 401

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RESULT 6

Q7T2T7 PRELIMINARY; PRT; 524 AA.

AC Q7T2T7;
 DT 01-OCT-2003 (TREMBLrel. 25, Created)
 DT 01-OCT-2003 (TREMBLrel. 25, Last sequence update)
 DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
 DE Methyl-cytosine binding protein 2.
 GN Name=Mecp2;
 OS Brachydanio rerio (Zebrafish) (Danio rerio).
 OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
 OC Cyprinidae; Danio.
 OK NCBI_TaxId=7955;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Coverdale L.E., Martin C.C.;
 RL Submitted (OCT-2003) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AY298900; AAP57248.2; -.
 DR HSSP: Q9UN29; 1IG4.
 DR ZFIN: ZDB-GENE-030131-7190; mec2.
 DR GO: GO:0003677; F:DNA binding; IEA.
 DR InterPro: IPR001739; Methyl-CpG_bind.
 DR Pfam: PF01429; MBD; 1.
 DR SMART: SM00391; MBD; 1.
 SQ SEQUENCE 524 AA; 57152 MW; B8593B4B84DCD1 CRC64;

Query Match 7.8%; Score 238.5; DB 2; Length 524;
 Best Local Similarity 24.4%; Pred. No. 2.2e-06;

Matches 106; Conservative 57; Mismatches 159; Indels 113; Gaps 18;

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QY 27 VPDPNDLRKEDVAMELERVEDEEQMMIKRSSCNPLQEPISAQGA---TAGTECR 83
DB 49 VPPPSLEFTORDVQQAQAE-AGKSE-----PI--DPEVQALSAESSASAKOR 93
QY 84 KSV-----PCGWERVVKORLFGTAGRFVYFISPGQLKFRSKSLANYLAK 130
DB 94 RSVLRDQPMTEBDSLPQWTRKLKQKSGSGSAGKFDVYLINPGKAFRSKVELMAYRQX 153
QY 131 NGESTLKPEDPDPFVYLSKRGIKSRVYKDCSMALTSHLQNGSNNSNMLRTSKCKQDVFM 190
DB 154 VGDITTDNDPDTFY-TQSGSPSR-----REKRPPKPKQV 188
QY 191 PPSSESLEQSRGLSNFTSHLLKEDBGVDVNFPRVKRKGKVTI----- 237
DB 189 KPS-----GRGRPRKSGKVRQAQTEGV--AVKRVLEKPGKLVMPVAPPTERGA 239
QY 238 -LKGIPIKTKKGGKSGFVQSDSKRESVYCNKADASEPVAQKSQLDRFVCI SDAGAC 296
DB 240 PLGQAPVAKARGRKR-----KSDPPSTPKRGRKRPATVSQS---TVGTSAAY 288
QY 297 GETLSVTSE-----ENSLVKKERSLSGSGNFCEQKTSGLINKFCSAKDEHNEKYE 349
DB 289 AAAAILTVAKKKALKESSAKPVQERALP-----IKKRTRETLIEL-EASTSATETFE 342
QY 350 DTFLESEIEIGKVEY-VERKEHLHTDLKRGSEMDNNSCPRKDTGTEKIQEDTIPRQ 408
DB 343 KRLTASTVTPTGESELETQCKPHKPS--RKIKHEADPGSSSGTTASG-----VAPKSH 393
QY 409 IERRKTSLYFSKYN 423
DB 394 KRRDQGRQFPHHMH 408

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RESULT 7

Q9YGC6 PRELIMINARY; PRT; 467 AA.

AC Q9YGC6;
 DT 01-MAY-1999 (TREMBLrel. 10, Created)
 DT 01-MAY-1999 (TREMBLrel. 10, Last sequence update)
 DT 05-JUL-2004 (TREMBLrel. 27, Last annotation update)
 DE Methyl-CpG-binding protein Mecp2 (Methyl-CpG-binding protein 2).
 GN Name=Mecp2;
 OS Xenopus laevis (African clawed frog).
 OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae; Pipidae;
 OC Xenopodinae; Xenopus.
 OK NCBI_TaxId=8355;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA MEDLINE=98282101; PubMed=9620779; DOI=10.1038/561;
 RX Jones P.L., Veenstra G.J.C., Wade P.A., Vermaak D., Kass S.U.,
 RA Landsberger N., Strouboulis J., Wolffe A.P.;
 RT "Methylated DNA and Mecp2 recruit histone deacetylase to repress
 transcription.";
 RT Nat. Genet. 19:187-191 (1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RA Kass S.U., Strouboulis J., Wolffe A.P.;
 RL Submitted (FEB-1998) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF106951; AAD03736.1; -.
 DR EMBL: AF051768; AAD02651.1; -.
 DR HSSP: P51608; 1OX9.
 DR GO: GO:0005634; C:nucleus; IEA.
 DR GO: GO:0003677; F:DNA binding; IEA.
 DR GO: GO:0003355; P:regulation of transcription; IEA.
 DR InterPro: IPR00637; A+T hook.
 DR InterPro: IPR001739; Methyl-CpG_bind.
 DR Pfam: PF02178; AT_hook; 1.
 DR Pfam: PF01429; MBD; 1.
 DR SMART: SM00384; AT hook; 2.
 DR SMART: SM00391; MBD; 1.

SQ SEQUENCE 467 AA; 51757 MW; 5D3A719A59E560BC CRC64;
Query Match 7.2%; Score 220.5; DB 2; Length 467;
Best Local Similarity 25.7%; Pred. No. 2.3e-05;
Matches 98; Conservative 60; Mismatches 157; Indels 67; Gaps 15;
OY 22 SSERLVPPNDLRKEDVAMELER---VGEDEEQWIMKRS--SECNPLQPIASAOQGA 76
DB 49 SSEH-QPEPADDEGADMSSEAEENTLAVPESSASPKGRSVIRDGPRYEDP----- 99
OY 77 TAGTECRKSVPCGMEVRRVVKRLFGKTAGRPDVYFISPOGLKFKRSKIANTLHKNGETSL 136
DB 100 -----TLPEGWTRKTKORRSGRSAGKFDVYLLINPGKAFRSKVELIAVFOKGDPTSL 151
OY 137 KPREDFTVLKRGKSKSYKXCSMAALTSHLQNGSNMNLFRTRSKCKOV--FMPSS 194
DB 152 DPNPDFTV-TGRGSPSRREQ-----KPKPKPKPKSVSGRGRGPKGSIKKVPPVK 204
OY 195 SSRLQESHRLNFTSTHLLK----EDEGVDDVFRKV-----RKPKGVTILKGP 242
DB 205 SEGQVAVKVIK-SFGKLVMPYSGTKEASDATTSQOVLYIKGGRKRGK-TPPSAP 262
OY 243 IKTTRKGRKSCSGFVQSDSRSEVYCNKADESEFVAKSGLDRTVCISDAGCGETLSV 302
DB 263 KKGKGRKPSNVSLAAMAAAKKAI---KESIKPLLE-----TVLPTRKRTRETTISV 313
OY 303 TSEE-----NSLVKK-----KERSLSGNSNFCSEKTSIGIINKPCASDSEHNEKE 349
DB 314 DVKDTIKPEPLTPYIEKWKMGQNPSPKSPSRSTEGSPKIKTGLPKKELQHHHHHHHHH 373
OY 350 DTFLSEERIGTKEVERKEHL 371
DB 374 HHSESKSATSPEPETSQNI 395
RESULT 8
O7SCQ2 PRELIMINARY; PRT; 682 AA.
AC O7SCQ2;
DT 01-MAR-2004 (TREMBlrel. 26, Created)
DT 01-MAR-2004 (TREMBlrel. 26, Last sequence update)
DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
DE Hypothetical protein.
GN Name=NCU09815.1;
OS Neurospora crassa.
OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;
OC Sordariomycetidae; Sordariales; Sordariaceae; Neurospora.
OX NCBI_TaxID=5141;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=OR74A;
RA Galagan J.E., Calvo S.E., Borkovich K.A., Selker E.U., Read N.D.,
RA Jaffe D., Fitzhugh W., Ma L.-D., Smirnov S., Purcell S., Reiman B.,
RA Elkins T., Engels R., Wang S., Nielsen C.B., Butler J., Endrizzi M.,
RA Qui D., Ianakiev P., Pedersen D., Nelson M., Washburne M.,
RA Selitrenikoff C.P., Kinsey J.A., Braun E.L., Zelter A., Schulte U.,
RA Koche G.O., Jedd G., Mewes W., Steben C., Marcotte E., Greenberg D.,
RA Roy A., Foley K., Naylor J., Thomann N., Barrett R., Gnetre S.,
RA Kamal M., Kanyseis W., Mancell E., Bielke C., Rudd S., Frisman D.,
RA Krystofova S., Rasmussen C., Metzberg R.L., Perkins D.D., Kroken S.,
RA Cogoni C., Macino G., Catchside D., Li W., Pratt R.J., Osmari S.A.,
RA Desouza C.C., Glass L., Orbach M.J., Berglund J., Voelker R.,
RA Yarden O., Plamann M., Seiler S., Dunlap J., Radford A., Aramayo R.,
RA Natvig D.O., Alex L.A., Mannhaupt G., Edbole D.J., Freitag M.,
RA Paulsen I., Sachs M.S., Lander E.S., Nussbaum C., Birren B.,
RA "The Genome Sequence of the Filamentous Fungus Neurospora crassa."
RT Nature 0:0-0(2003).
CC -!- CAUTION: The sequence shown here is derived from an
EMBL/GenBank/DBJ whole genome shotgun (WGS) entry which is
preliminary data.
CC EMBL; AABX01000087; EAA34519.1; -!
DR GO; GO:0006284; P:base-excision repair; IEA.
DR InterPro; IPR011257; DNA_glycylase.

DR InterPro; IPR003265; Endo_3c.
DR Pfam; PF00730; Hnh-GPD; 1.
KW Hypothetical protein.
SQ SEQUENCE 662 AA; 75928 MW; 4PDCCAA26102EB84 CRC64;
Query Match 7.0%; Score 213; DB 2; Length 682;
Best Local Similarity 20.8%; Pred. No. 0.00011;
Matches 93; Conservative 62; Mismatches 155; Indels 138; Gaps 14;
OY 240 GIPIKTKKGRKSCSGF-VQSDSKRS-----VNKKADESEFVAKSGL 284
DB 164 GKRVKTKTGWGRSVFMAODDPKTSROYTRGRGETGVSGAVRKAMNGHEPISGCK- 222
OY 285 DRTVCISDAGCGETLVTSENSLVKKK-RLSLSGSNFCSEKTSIGIINKPCASDSE 343
DB 223 -----MLPFRWNSVVRSAPOGGBE---SKAKSRLLDK-TKLKHSK 263
OY 344 HNEK-YEDTFLESEEIGTKEVERKEHLHTDLKXGSEMDDNC-SPTKDFTEKIFQ 400
DB 264 FPDEPSYADTWSPQSLASHSSPNEMLDVALGVSVAGTSSPDECVLAAMVRIETERTYK 323
OY 401 EDTI---PRQIERKTSLYFSSKYKNEALSPRRKAFKKTTP-----RSPFN 446
DB 324 SPFSEDPMPPISTSKASPPKSTKTSKRSPTKKS--RNHPRGISCLPIAPISARFG 381
OY 447 LVQETLPHDPMKLLIATIFLNRTSGKMAIPVLMFKLEKYSAEVARTADWDSLEIKPL 506
DB 382 LIQEVAAADPFRLLIANTFLIKRGMTAIPFLRQMDLFTPLPRLASADSEIINLRPL 441
OY 507 GLYDLRAKTIKVS-----DEYLTQWKYPIELHGI----- 537
DB 442 GLSVNRCSVLQKVARMPFIECPCKEKRYGVKNYPRPDAGVAGKQGFTEGPDFFHIGKA 501
OY 538 -----GKYGNSYRIFCN----- 551
DB 502 SQAEFDDDDDRINAIRKAKERHAIGAMEIGHLTGQFPALDSWRIFCRDKLLGRADWKGK 561
OY 552 -----EMKQVPEBDKINKYHDMWL 571
DB 562 GRHPEPQPEMWRVLPQDKELRAYLRMMW 589
RESULT 9
MEC2 HUMAN
ID MEC2 STANDARD; PRT; 486 AA.
AC P51608; O15233;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Methyl-CpG-binding protein 2 (MecP-2 protein) (MecP2).
GN Name=MecP2;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Kudo S., Fukuda M.,
RL Submitted (SSP-1995) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RA Thiesen J., Straetling W.H.,
RL Submitted (APR-1997) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE=97130625; PubMed=9976388;
RA Vilain A., Apioy F., Vogt N., Dutrillaux B., Malfoy B.,
RT "Assignment of the gene for methyl-CpG-binding protein 2 (MecP2) to
human chromosome band Xq28 by in situ hybridization."
RL Cytogenet. Cell Genet. 74:293-294 (1996).
RN [4]
RP SEQUENCE FROM N.A.

RA Reichwald K., Rosenthal A., Kioschis P., Platzter M.,
 RT "Mapping and sequence analysis of the human MECP2 locus.",
 RL Submitted (Oct-1997) to the EMBL/GenBank/DBJ databases.
 [5]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=9929240; PubMed=10369871; DOI=10.1093/hmg/8.7.1253;
 RA Coy J.F., Sedlacek Z., Baechner D., Delius H., Poustka A.,
 RT "A complex pattern of evolutionary conservation and alternative
 RT polyadenylation within the long 3'-untranslated region of the methyl-
 RT Cpg-binding protein 2 gene (MECP2) suggests a regulatory role in gene
 RT expression.";
 RL Hum. Mol. Genet. 8:1253-1262 (1999).
 [6]
 RP SEQUENCE FROM N.A.
 RA Reichwald K., Thiesen J., Wiehe T., Weitzel J., Straetling W.H.,
 RA Kioschis P., Rosenthal A., Platzter M.,
 RT "Comparative sequence analysis of the MECP2-1 locus in human and mouse
 RT reveals new untranslated regions";
 RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
 [7]
 RP SEQUENCE FROM N.A.
 RC TISSUE=placenta;
 RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
 RA Struhsberg R.L., Reingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shemen C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhac N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diachenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stalenon M., Soares M.B., Bonaldi M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Usdin T.B., Toshlyuk S., Carninci P., Prange C.,
 RA Rana S.S., Loquellano N.A., Peters G.J., Abtenson R.D., Mallary S.J.,
 RA Bosak S.A., Mesman P.J., McKernan K.U., Malek J.A., Gunaratne P.H.,
 RA Richards S., Morley K.C., Hale S., Garcia A.M., Gay L.J., Hulys S.W.,
 RA Villalón D.K., Murry D.W., Sodergren B.J., Lu X., Gibbs R.A.,
 RA Fahy J., Helton E., Ketterman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.W.,
 RA Buterfield V.S.N., Krzywinski M.I., Skalska U., Smalins D.E.,
 RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.,
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
 [8]
 RP SEQUENCE OF 10-486 FROM N.A.
 RC TISSUE=skeletal muscle;
 RX MEDLINE=96327611; PubMed=8672133;
 RA D'Esposito M., Quadert N.A., Ciccodicola A., Bruni P., Esposito T.,
 RA D'Urbio M., Brown S.D.M.,
 RT "Isolation, physical mapping, and Northern analysis of the X-linked
 RT human gene encoding methyl Cpg-binding protein, MECP2.";
 RL Mamm. Genome 7:533-535 (1996).
 [9]
 RP SEQUENCE OF 10-486 FROM N.A.
 RA Reichwald K., Bauer D., Brenner V., Drescher B., Coy J.F.,
 RA Kioschis P., Korn B., Myakatura G., Platzter M., Poustka A.,
 RA Sandoval N., Rosenthal A.,
 RT Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
 [10]
 RP REVIEW ON VARIANTS.
 RX PubMed=12872250; DOI=10.1002/humu.10243;
 RA Miltenberger-Miltenyi G., Laccone F.,
 RT "Mutations and polymorphisms in the human methyl Cpg-binding protein
 RT MECP2.";
 RL Hum. Mutat. 22:107-115 (2003).
 [11]
 RP VARIANTS RTT TRP-106; CYS-133; SER-155; MET-158 AND CYS-306, AND
 RP VARIANT LYS-397.
 RX PubMed=10577905;
 RA Wan M., Lee S.S.J., Zhang X., Houwink-Manville I., Song H.-R.,
 RA Amir R.E., Budden S., Naidu S., Pereira J.L.P., Lo I.F.M.,
 RA Zoghbi H.Y., Schanen N.C., Francke U.,
 RT "Ret syndrome and beyond: recurrent spontaneous and familial MECP2

RT mutations at Cpg hotspots.";
 RL Am. J. Hum. Genet. 65:1520-1529 (1999).
 [12]
 RP VARIANTS RTT TRP-106; CYS-133; SER-155 AND MET-158.
 RX MEDLINE=99438392; PubMed=10508514; DOI=10.1038/13810;
 RA Amir R.E., Van den Veyver I.B., Wan M., Tiran C.Q., Francke U.,
 RA Zoghbi H.Y.,
 RT "Ret syndrome is caused by mutations in X-linked MECP2, encoding
 RT methyl-Cpg-binding protein 2.";
 RL Nat. Genet. 23:185-188 (1999).
 [13]
 RP INVOLVEMENT IN X-LINKED MENTAL RETARDATION WITH PROGRESSIVE
 RP SPASTICITY.
 RX PubMed=10986043;
 RA Meloni I., Brutini M., Longo I., Mari F., Rizzolio F., D'Adamo P.,
 RA Denvirliend K., Fryns J.-P., Tonello D., Renieri A.,
 RT "A mutation in the Ret syndrome gene, MECP2, causes X-linked mental
 RT retardation and progressive spasticity in males.";
 RL Am. J. Hum. Genet. 67:982-985 (2000).
 [14]
 RP VARIANTS RTT VAL-100; GLN-106; TRP-106; CYS-133; ARG-152; SER-155;
 RP MET-158; ARG-305; CYS-306 AND HIS-306, AND VARIANTS CYS-86; MET-203;
 RP PRO-287; ALA-291; LYS-397; ILE-412 AND THR-444.
 RX PubMed=11055898;
 RA Buysse I.M., Pang P., Hoon K.T., Amir R.E., Zoghbi H.Y., Roa B.B.,
 RT "Diagnostic testing for Ret syndrome by DHPLC and direct sequencing
 RT analysis of the MECP2 gene: identification of several novel mutations
 RT and polymorphisms.";
 RL Am. J. Hum. Genet. 67:1428-1436 (2000).
 [15]
 RP VARIANT MEX16 VAL-140, AND VARIANT MET-203.
 RX MEDLINE=20465115; PubMed=11007980; DOI=10.1016/S0014-5793(00)01994-3;
 RA Orrico A., Lam C., Gall L., Dotti M.T., Hayek G., Tong S.F.,
 RA Poon P.M., Zappella M., Federico A., Sorrentino V.,
 RT "MECP2 mutation in male patients with non-specific X-linked mental
 RT retardation.";
 RL FEBS Lett. 481:285-288 (2000).
 [16]
 RP VARIANTS RTT LEU-101; HIS-101; THR-101; TRP-106; CYS-133; CYS-134;
 RP ARG-152; MET-158; ARG-225; LEU-302; CYS-306 AND HIS-306, AND VARIANTS
 RP LEU-229 AND THR-439.
 RX PubMed=10767337; DOI=10.1093/hmg/9.7.1119;
 RA Chesdale J.P., Gill H., Fleming N., Maynard J., Kerr A., Leonard H.,
 RA Kravczak M., Cooper D.N., Lynch S., Thomas N., Hughes H., Hulten M.,
 RA Ravine D., Sampson J.R., Clarke A.,
 RT "Long-read sequence analysis of the MECP2 gene in Ret syndrome
 RT patients: correlation of disease severity with mutation type and
 RT location.";
 RL Hum. Mol. Genet. 9:1119-1129 (2000).
 [17]
 RP VARIANTS RTT GLN-106; MET-158; ARG-302; CYS-306 AND ALA-322.
 RX PubMed=10814719; DOI=10.1093/hmg/9.9.1377;
 RA Bienvenu T., Carrie A., de Roux N., Vinet M.-C., Jonveaux P.,
 RA Couvert P., Villard L., Arizmanoglu A., Beidjord C., Fontes M.,
 RA Tardieu M., Chelly J.,
 RT "MECP2 mutations account for most cases of typical forms of Ret
 RT syndrome.";
 RL Hum. Mol. Genet. 9:1377-1384 (2000).
 [18]
 RP VARIANTS RTT MET-158; HIS-302 AND CYS-306, AND VARIANTS VAL-201;
 RP ALA-232; LEU-251 AND SER-376.
 RX PubMed=10944854;
 RA Amano K., Nomura Y., Segawa M., Yamakawa K.,
 RT "Mutational analysis of the MECP2 gene in Japanese patients with Ret
 RT syndrome.";
 RL J. Hum. Genet. 45:231-236 (2000).
 [19]
 RP VARIANTS RTT TRP-106; PHE-124; CYS-133; CYS-134; ARG-152; MET-158 AND
 RP CYS-306.
 RX MEDLINE=20439334; PubMed=10991688;
 RA Obata K., Matsumi T., Yamashita Y., Fukuda T., Kuwajima K.,
 RA Horiuchi I., Nagamitsu S., Iwanaga R., Kimura A., Omori I., Endo S.,
 RA Mori K., Kondo I.,

"Mutation analysis of the methyl-CpG binding protein 2 gene (MECP2) in RT patients with Rett syndrome.";
RT J. Med. Genet. 37:608-610(2000).
RN [20]
RP VARIANTS RTT ARG-101; TRP-106; MET-158 AND CYS-306, AND VARIANT RP LYS-397.
RX MEDLINE=20439335; PubMed=10991689;
RA Hampson K., Woods C.G., Jaffif F., Webb T.;
RT "Mutations in the MECP2 gene in a cohort of girls with Rett syndrome.";
RL J. Med. Genet. 37:610-612(2000).
RN [21]
RP VARIANT RTT HIS-133.
RX PubMed=11706982; DOI=10.1002/ana.1272;
RA Armstrong J., Poo P., Pineta M., Albar E., Gean E., Catala V.,
RT "Classic Rett syndrome in a boy as a result of somatic mosaicism for a MECP2 mutation.";
RL Ann. Neurol. 50:692-692(2001).
RN [22]
RP VARIANTS RTT TRP-106; CYS-134; ARG-152; MET-158; ALA-302; CYS-306 AND RP ALA-322, AND VARIANTS VAL-201 AND LYS-397.
RX PubMed=11738883; DOI=10.1016/S0387-7604(01)00342-4;
RA Giunti L., Pelagatti S., Lazzerini V., Guarducci S., Lapi E.,
RA Covello S., Cecconi A., Ombroni L., Andreucci E., Sami I.,
Query Match 6.8%; Score 208.5; DB 1; Length 486;
Best Local Similarity 23.9%; Pred. No. 0.00013;
Matches 104; Conservative 53; Mismatches 149; Indels 129; Gaps 17;
QY 37 EDVAMELERVGEDEQOMIKRSSSECNPLLOEPIASAQFG-----ATAGT 80
DB 22 KDXPLKFKKKVKKDKKEKEGHEPVQSAHNSABPEAGKAFETSEGSAPAVPEASASP 81
QY 81 ECRKSV-----PCGWERVVKORLFGKTAGRPDYVFISPGGLKFRSKSLANY 127
DB 82 KQRSIIIRDGRPMYDDPTLPEGWTRKDKQRKSGRSAGKYDYLINPQKAFRSKVELIAY 141
QY 128 LHKNGETSLKPEDPFTVLSKRGTKSRKYKDCSMAALTSHLQNSNNSNMLRTRSKCKD 187
DB 142 FEKVGDTSLDPNDFPTV-TGRGSPSR-----REQPPPK- 175
QY 188 VFMPSSSSSELQESRGL---SNFTSTHLLEKDEGVDDVNFKVRKPKGKVTILKGIPIK 244
DB 176 ----PKSPKAPGTGRGRPRKSGGTTTRPKATSEGVQK--RVLEKSPGK--LIVKMPF- 226
QY 245 KTKKGCRKSCSGFVQSDS---KRESVCNKADAESEPVAKXSQLDRTVCISDAGACETL 300
DB 227 QTSFGKAEAGGATTSQVWVVKRPGKRKAADPQALPKKR-----GRKPGSVV 276
QY 301 SVTSEENSLVKKERSLSSGSNFCSEQKTSGLINKFCASDSEHNEKYEFTLSESEIGT 360
DB 277 AAAAEEAKKAVKSSIR-----SVQETVLPIKK-----RKTRET-----V 312
QY 361 KVEVVERKEHLHTDIL--KRGSEMDNNSPTRKDFTEGKIFQEDTIPRTQIERKTSLYF 418
DB 313 STEVEVVKPLVSTLGEKSGKGLTKCKSPGRK-----SKESPGR----- 354
QY 419 SSKYKKEALSPPRK 433
DB 355 ----SSSASSPKKE 365
RESULT 10
Q6QH9 PRELIMINARY; PRT; 498 AA.
AC Q6QH9;
DT 05-JUL-2004 (TREMBLrel. 27, Created)
DT 05-JUL-2004 (TREMBLrel. 27, Last sequence update)
DT 05-JUL-2004 (TREMBLrel. 27, Last annotation update)
DE Methyl CpG binding protein 2 isoform B.
GN Name=MECP2;
OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX PubMed=15034579; DOI=10.1038/ng1327;
RA Matczakian G.N., Lohi H., Munteanu I., Alfred S.E., Yamada T.,
RA MacLeod P.J., Jones J.R., Scherer S.W., Schanen N.C., Friez M.J.,
RA Vincent J.B., Mnasasian B.A.;
RT "A previously unidentified MECP2 open reading frame defines a new protein isoform relevant to Rett syndrome.";
RL Nat. Genet. 36:339-341(2004).
DR EMBL; AY541280; AAS5455.1; -.
DR GO; GO:0005634; C:nucleus; IEA.
DR GO; GO:0003677; F:DNA binding; IEA.
DR GO; GO:0006355; P:regulation of transcription, DNA-dependent; IEA.
DR InterPro; IPR000637; A+T_hook.
DR InterPro; IPR001739; Methyl-CpG_bind.
DR Pfam; PF02178; AT_hook; 1.
DR Pfam; PF01429; MBD; 1.
DR SMART; SM00391; MBD; 1.
SQ SEQUENCE 498 AA; 53323 MW; 443BGB3D5EA4DAB8 CRC64;
Query Match 6.8%; Score 208.5; DB 2; Length 498;
Best Local Similarity 23.9%; Pred. No. 0.00014;
Matches 104; Conservative 53; Mismatches 149; Indels 129; Gaps 17;
QY 37 EDVAMELERVGEDEQOMIKRSSSECNPLLOEPIASAQFG-----ATAGT 80
DB 34 KDXPLKFKKKVKKDKKEKEGHEPVQSAHNSABPEAGKAFETSEGSAPAVPEASASP 93
QY 81 ECRKSV-----PCGWERVVKORLFGKTAGRPDYVFISPGGLKFRSKSLANY 127
DB 94 KQRSIIIRDGRPMYDDPTLPEGWTRKDKQRKSGRSAGKYDYLINPQKAFRSKVELIAY 153
QY 128 LHKNGETSLKPEDPFTVLSKRGTKSRKYKDCSMAALTSHLQNSNNSNMLRTRSKCKD 187
DB 154 FEKVGDTSLDPNDFPTV-TGRGSPSR-----REQPPPK- 187
QY 188 VFMPSSSSSELQESRGL---SNFTSTHLLEKDEGVDDVNFKVRKPKGKVTILKGIPIK 244
DB 188 ----PKSPKAPGTGRGRPRKSGGTTTRPKATSEGVQK--RVLEKSPGK--LIVKMPF- 238
QY 245 KTKKGCRKSCSGFVQSDS---KRESVCNKADAESEPVAKXSQLDRTVCISDAGACETL 300
DB 239 QTSFGKAEAGGATTSQVWVVKRPGKRKAADPQALPKKR-----GRKPGSVV 288
QY 301 SVTSEENSLVKKERSLSSGSNFCSEQKTSGLINKFCASDSEHNEKYEFTLSESEIGT 360
DB 289 AAAAEEAKKAVKSSIR-----SVQETVLPIKK-----RKTRET-----V 324
QY 361 KVEVVERKEHLHTDIL--KRGSEMDNNSPTRKDFTEGKIFQEDTIPRTQIERKTSLYF 418
DB 325 STEVEVVKPLVSTLGEKSGKGLTKCKSPGRK-----SKESPGR----- 366
QY 419 SSKYKKEALSPPRK 433
DB 367 ----SSSASSPKKE 377
RESULT 11
Q72384 PRELIMINARY; PRT; 516 AA.
AC Q72384;
DT 01-OCT-2003 (TREMBLrel. 25, Created)
DT 01-OCT-2003 (TREMBLrel. 25, Last sequence update)
DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
DE Hypothetical protein DKFZp686A24160 (Fragment).
GN Name=DKFZp686A24160;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;


```

[1]
RN SEQUENCE FROM N.A.
RC TISSUE=human colon endothelial primary cell culture;
RA Bloeker H., Boecker M., Mewes H.W., Well B., Amid C., Osanger A.,
RA Robo G., Han M., Wiemann S.;
RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL: BX538060; CAD97991.1; -.
DR HSP; Q9UN29; 1164.
DR GO: GO:0005634; C:nucleus; IEA.
DR GO: GO:0003677; F:DNA binding; IEA.
DR GO: GO:0003355; P:regulation of transcription, DNA-dependent; IEA.
DR InterPro: IPR000637; A+T hook.
DR InterPro: IPR001739; Methyl-Cpg_bind.
DR Pfam: PF02178; AT hook; 1.
DR Pfam: PF01429; MBD; 1.
DR SMART: SM00384; AT hook; 2.
DR SMART: SM00391; MBD; 1.
DR Hypothetical protein.
KW NON_TER
SQ SEQUENCE 516 AA; 55204 MW; 27CD37B9164176B0 CRC64;

Query Match
Best Local Similarity 23.9%; Score 208.5; DB 2; Length 516;
Matches 104; Conservative 53; Mismatches 149; Indels 129; Gaps 17;

QY 37 EDVAMELERVGEDEBQMIKRSSECNPLLOEPASAOFG-----ATAGT 80
DB 52 KDKPLKFKVKKDKKEKKGKHEPVQPSAHNSAEPABAGKAETSEGSAGAPAVEASASP 111
QY 81 ECRKSV-----PCGWERVVKQRLFGKTAGRDVYFISQGLKPKSKSLANY 127
DB 112 KQRRSIIIRDGPWYDPTLPBEGWTRKTKQKRSKRSAGKYDVLINPQGAFFSKVELLAY 171
QY 128 LKNGETSLKPEPDFTVLSKRGISRYDCSMALTSHLQNSNNMNLRTSRCKKD 187
DB 172 FEKVGDTSLDPNDPFTV-TGRGSPR-----REQRPKK- 205
QY 188 VEMPSSSELOESRGL---SNFTSTHLLKEDGVDDVNFRRKVRKPKGKVTILKGIPIK 244
DB 206 ----PSPKAPGTGGRGRPKSGCTTRPRAATSEGVQK--RVLEKSPK--LIVMP- 256
QY 245 KTKKCGRKSCKSGFVQSDS---KRSVCKKADABEPVAKQSLDRTVCISDAGAGETL 300
DB 257 QTSFGCKAGGAGATTSTQVMVIRPKRKRAEADPQAIPIKPKR-----GRKGSYV 306
QY 301 SVTSEENSLVKKKERSLSSGNSFCSEOKTSGIINFKCSAKDSEHNEKYEDTFLSEBEIGT 360
DB 307 AAAAABAKKAAKAVKSSIR-----SVQETVLPKK-----RKTRET-----V 342
QY 361 KVEVERKEHLHTDIL--KRGSEMDNNSCFTRKDTGEGKIFQEDTTPRTQIERRTSLYF 418
DB 343 STEVEKVVPLVSTLGEKSGKGLTKCKSPGR-----SKESSPKGR----- 384
QY 419 SKRTKEALSPRRK 433
DB 385 ----SSASASPCKE 395

RESULT 12
MEC2_MACFA STANDARD; PRT; 486 AA.
AC 0951G8;
DT 25-OCT-2004 (Rel. 45, Created)
DT 25-OCT-2004 (Rel. 45, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Methyl-Cpg-binding protein 2 (Mecp-2 protein) (Mecp2).
GN Name=Mecp2;
OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).
OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
OC Cercopithecinae; Macaca.
OC NCBL_taxonomy=9541;
RN [1]

```

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RP SEQUENCE FROM N.A.
RA Muramatsu S.;
RT "Excessive hand-wringing in a MPP-treated monkey.";
RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
CC -! FUNCTION: Chromosomal protein that binds to methylated DNA. It can
CC bind specifically to a single methyl-Cpg pair. It is not
CC influenced by sequences flanking the methyl-Cpgs. Mediates
CC transcriptional repression through interaction with histone
CC deacetylase and the corepressor SIN3A.
CC -! SUBUNIT: Interacts with FMBP3 (by similarity).
CC -! SUBCELLULAR LOCATION: Nuclear. Colocalized with methyl-Cpg in the
CC genome (by similarity).
CC -! SIMILARITY: Contains 2 A.T hook DNA-binding repeats.
CC -! SIMILARITY: Contains 1 methyl-Cpg-binding (MBD) domain.
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: AF295597; AAK97131.1; -.
DR HSP; P51608; 10K9.
DR InterPro: IPR000637; AT hook.
DR InterPro: IPR001739; Methyl-Cpg_bind.
DR Pfam: PF02178; AT hook; 1.
DR Pfam: PF01429; MBD; 1.
DR SMART: SM00384; AT hook; 2.
DR SMART: SM00391; MBD; 1.
DR PROSITE: PS50382; MBD; 1.
DR DNA-binding; Nuclear protein; Repeat; Repressor;
KW Transcription regulation.
FT DOMAIN 90 162 MBD.
FT DNA_BIND 185 197 A.T hook 1.
FT DNA_BIND 265 277 A.T hook 2.
FT DOMAIN 366 372 His-rich.
FT DOMAIN 405 405 Pro-rich.
SQ SEQUENCE 486 AA; 52426 MW; 3471B61D90D92A7D CRC64;

Query Match
Best Local Similarity 23.9%; Score 207.5; DB 1; Length 486;
Matches 104; Conservative 53; Mismatches 149; Indels 129; Gaps 17;

QY 37 EDVAMELERVGEDEBQMIKRSSECNPLLOEPASAOFG-----ATAGT 80
DB 22 KDKPLKFKVKKDKKEKKGKHEPVQPSAHNSAEPABAGKAETSEGSAGAPAVEASASP 81
QY 81 ECRKSV-----PCGWERVVKQRLFGKTAGRDVYFISQGLKPKSKSLANY 127
DB 82 KQRRSIIIRDGPWYDPTLPBEGWTRKTKQKRSKRSAGKYDVLINPQGAFFSKVELLAY 141
QY 128 LKNGETSLKPEPDFTVLSKRGISRYDCSMALTSHLQNSNNMNLRTSRCKKD 187
DB 142 FEKVGDTSLDPNDPFTV-TGRGSPR-----REQRPKK- 175
QY 188 VEMPSSSELOESRGL---SNFTSTHLLKEDGVDDVNFRRKVRKPKGKVTILKGIPIK 244
DB 176 ----PSPKAPGTGGRGRPKSGCTTRPRAATSEGVQK--RVLEKSPK--LIVMP- 226
QY 245 KTKKCGRKSCKSGFVQSDS---KRSVCKKADABEPVAKQSLDRTVCISDAGAGETL 300
DB 227 QTSFGCKAGGAGATTSTQVMVIRPKRKRAEADPQAIPIKPKR-----GRKGSYV 276
QY 301 SVTSEENSLVKKKERSLSSGNSFCSEOKTSGIINFKCSAKDSEHNEKYEDTFLSEBEIGT 360
DB 277 AAAAABAKKAAKAVKSSIR-----SVQETVLPKK-----RKTRET-----V 312
QY 361 KVEVERKEHLHTDIL--KRGSEMDNNSCFTRKDTGEGKIFQEDTTPRTQIERRTSLYF 418
DB 343 STEVEKVVPLVSTLGEKSGKGLTKCKSPGR-----SKESSPKGR----- 354

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```
QY      419 SSKYNKEALSPRRK 433
          : | | | | :
Db      355 ----SSSASSPKKE 365
```

RESULT 13	
042403	
ID 042403	PRELIMINARY; PRT; 344 AA

RT "Chicken MAR-binding protein ARBP is homologous to rat methyl-CpG
RT binding protein MeCP2.";
RT Mol. Cell. Biol. 17:5656-5666 (1997).
RL

RT "Comparative sequence analysis of the MEC2-locus in human and mouse
RT reveals new transcribed regions.";
RL Mamm. Genome 11:182-190(2000).

Query Match	6.6%	Score	202.5	DB 2	Length	344			
Best Local Similarity	34.7%	Pred. No.	0.0002						
Matches	59	Conservative	17	Mismatches	57	Indels	37	Gaps	5

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QY      16 AAFVTSSSELTVPDPNDUKEDVNAELER-----VGBDEBDMTKRSECNPLLOEIA 70
Db      5  AAAAAGEEEL-----EEQADBSVALKMRPPRAKKGREREDPEALAEAPGAPAE 59
QY      71 SAQGFATAGTEC-----RKSV-----PCGMEVAVQRLFGKTAG 104
Db      60 AGKDGSGGTAAPAVPEASAPSKORSTIRDRGPNYDPTLPSBMTKTLKORSGRSAG 119
QY      105 RFVDYFISPGQLFKRSKSSILANYLHNQGETSLKPEDPDTVLAKRGIKSR 154
Db      120 KYDVIYLNPGQKAFRSKVELIAFEKVGQTSLDPNDFPTV--TGGSFSR 168

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RESULT 14		
MEC2_RAT		
ID	MEC2_RAT	STANDARD; PRT; 492 AA
AC	Q00566;	
DT	01-APR-1993	(Rel. 25, Created)
DT	01-APR-1993	(Rel. 25, Last sequence update)
DT	25-OCT-2004	(Rel. 45, Last amortation update)

DE Mechyl_Cpg-binding protein 2 (MeCP-2 protein) (MECP2).
GN Name=MeCP2;
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Rattus
OX NCBI_TaxId=10116;

CC
CC -I- FUNCTION: Chromosomal protein that binds to methylated DNA. It can
CC bind specifically to a single methyl-CpG pair. It is not
CC influenced by sequences flanking the methyl-CpGs. Mediates
CC transcriptional repression through interaction with histone
CC deacetylase and the corepressor SIN3A (By similarity).
CC -SUBUNIT: Interacts with PMP3 (By similarity).
CC -SUBCELLULAR LOCATION: Nuclear. Colocalized with methyl-CpG in the
CC genome.
CC
CC -TISSUE SPECIFICITY: Present in all adult somatic tissues tested.
CC
CC -SIMILARITY: Contains 2 A-T hook DNA-binding repeats.
CC
CC -SIMILARITY: Contains 1 methyl-CpG-binding (MBD) domain.
CC
CC -----
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CC or send an email to license@ebi.ac.uk).
CC

Query Match	6.6%	Score 201	DB 1	Length 492
Best Local Similarity	22.9%	Pred. No. 0.00039		
Matches 108; Conservative	57	Mismatches 156	Indels 150	Gaps 18

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QY 34 LRKE-----DVAMELERGCEDEGOMMIRKSSCNPLLQEPINSAQFG----- 75
    |||
Db 8 LRKESBEDDILQGLKEKPLKFKRKVKQDKDEKEGHEPLQPSAHNSAEPALGAETSES 67
    |||
QY 76 -----ATAGTECRKSV-----PCGMERVVKORLFGTAGRFVYFISP 113
    |||
Db 68 SGSAFAVFAASAPKORRSITRDRGPMWDDPTLDEGWTBKLRKQKSKSGSAGKIVYTLNP 127
    |||
QY 114 QGLKRSKSSILANYLHKNGESTLKPEDPFTVLRSKRGIKSYKDCOSMAALTSHLONSGN 173
    |||
Db 128 QGKAFRSKVELIATFEKKGDTSLDNPDPFLV-IGRGSPSK----- 167
    |||
QY 174 SNMNLRTSRKCKKQVMPSSSSSELQBSRGSLNFTSTHLLLKEDBGVDVNFRAV-RKPK 232
    |||
Db 168 -----REQKPKPK-----PKSPKPAKPTGTGRGGRPKSGSTGRPKAASGAVGVNKLTEKSP 217
    |||

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QY 233 GKVTLKGIPIKTKKGGKSCSGFVQSDS-----KRESVCKNKADESEPVNAQKSDRT 287
 DB 218 GK--LLVKKPFOASPGG--KGEGGATTSAGVWIKRPPKRAKADAPQALPKR----- 268
 QY 288 VQISNAGAGETLSTSENSLVKKKERSLSGNSFCSPKQTSGLINKFCSAKDBHNK 347
 DB 269 -----GRRPGSVVAAAAAARAKKAVKESIR-----SVQETVLPIK-----RK 307
 QY 348 YEDTFLESEIGTKYEVVERKEHLTDIL--KRSEMDNKNCSPTKEDFTGEKIFQEDTIP 405
 DB 308 TRET-----VSLFEVAVVPRPLVSTLTGKSGKGLTKCKSPGRK-----SKESP 351
 QY 406 RTQIERRRTSLYFSKKNKALSPRR-----KAFKMTPPRSP 444
 DB 352 KGR-----SSSASSPPKKEHHHHHAESPKAQMPPLPPPP 388
 RESULT 15
 ID MEC2 MOUSE STANDARD; PRT; 484 AA.
 AC 0922D6;
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 25-OCT-2004 (Rel. 45, Last annotation update)
 DT Methyl-CpG-binding protein 2 (Mecp2 protein) (Mecp2).
 GN Name=Mecp2;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 NC NCB1_TaxID=10090;
 RP [1]
 RN SEQUENCE FROM N.A.
 RA STRAIN=CS7B1/6;
 RC MEDLINE=98449942; PubMed=9774669;
 RX Hendrich B., Bird A.;
 RT "Identification and characterization of a family of mammalian methyl-
 CpG binding proteins.";
 RL Mol. Cell. Biol. 18:6538-6547(1998).
 RN [2]
 RN SEQUENCE FROM N.A.
 RA MEDLINE=99299240; PubMed=10369871; DOI=10.1093/hmg/8.7.1253;
 RC Coy J.F., Sedlacek Z., Beecher D., Delius H., Founteka A.;
 RX "A complex pattern of evolutionary conservation and alternative
 polyadenylation within the long 3'-untranslated region of the methyl-
 CpG-binding protein 2 gene (Mecp2) suggests a regulatory role in gene
 expression.";
 RL Hum. Mol. Genet. 8:1253-1262(1999).
 RN [3]
 RN SEQUENCE FROM N.A.
 RA Reichwald K., Thiesen J., Wiehe T., Kloeckle P., Straetling W.H.,
 RA Rosenthal A., Platzer M.;
 RT "Comparative analysis of the methyl CpG binding protein 2 locus in man
 and mouse reveals new untranslated sequences.";
 RT Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RN SEQUENCE FROM N.A.
 RA STRAIN=FVB/N; TISSUE=Mammary gland;
 RC MEDLINE=22386257; PubMed=12477932; DOI=10.1073/pnas.242603899;
 RX Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altschul S.F., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Ueda N.T., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Muljani S.J.,
 RA Bosak S.A., McMan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hultk S.W.,
 RA Vallion D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Rahney J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakeley R.W., Touchman J.W., Green E.D., Dickson M.C.,

RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield V.S.N., Krzywinski M.I., Skalske U., Smalhus D.E.,
 RA Schermer A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [5]
 RP INTERACTION WITH FMBP3.
 RX MEDLINE=97315177; PubMed=9171351; DOI=10.1093/emboj/16.9.2376;
 RA Bedford M.T., Chan D.C., Leder P.;
 RT "FMBP WW domains and the Abi SH3 domain bind to a specific class of
 RT proline-rich ligands.";
 RL EMBO J. 16:2376-2383(1997).
 CC -1- FUNCTION: Chromosomal protein that binds to methylated DNA. It can
 CC bind specifically to a single methyl-CpG pair. It is not
 CC influenced by sequences flanking the methyl-CpGs. Mediates
 CC transcriptional repression through interaction with histone
 CC deacetylase and the corepressor Sin3A (By similarity).
 CC -1- SUBUNIT: Interacts with FMBP3.
 CC -1- SUBCELLULAR LOCATION: Nuclear. Colocalized with methyl-CpG in the
 CC genome.
 CC -1- SIMILARITY: Contains 2 A.T hook DNA-binding repeats.
 CC -1- SIMILARITY: Contains 1 methyl-CpG-binding (MBD) domain.
 CC -----
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CC EMBL; AF072251; AAC68880.1; -;
 DR EMBL; AJ123922; CAB46495.1; -;
 DR EMBL; AF121351; -; NOT ANNOTATED_CDS.
 DR EMBL; AF158181; AAF33024.1; -;
 DR EMBL; BC027153; AAH27153.1; -;
 DR HSPF; P51608; IOK9.
 DR MGD; MGI:99918; Mecp2.
 DR GO; GO:0005634; C:nucleus; IDA.
 DR InterPro; IPR000637; A+T hook.
 DR InterPro; IPR001739; Methyl-CpG_bind.
 DR Pfam; PF01429; MBD; 1.
 DR SMART; SM00384; AT_hook; 2.
 DR PROSITE; PS50982; MBD; 1.
 DR DNA-binding; Nuclear protein; Repeat; Repressor;
 KW Transcription regulation.
 FT DOMAIN 90 162 MBD.
 FT DNA_BIND 185 197 A.T hook 1 (By similarity).
 FT DNA_BIND 265 277 A.T hook 2 (By similarity).
 FT DOMAIN 366 372 His-rich.
 FT FT DOMAIN 379 403 Pro-rich.
 SQ SEQUENCE 484 AA; 52307 MW; 62FD228F0118A49F CRC64;

Query Match 6.5%; Score 198.5; DB 1; Length 484;
 Best Local Similarity 24.1%; Pred. No. 0.00054;
 Matches 91; Conservative 50; Mismatches 154; Indels 83; Gaps 11;

QY 76 ATAGTECRKSV-----PCGMEVVYKORLFGKTAGRPDYVFTSPQGLKFRSKS 122
 DB 77 AASAPRQSRSTRDNGPWDTPLDEGWTRKLOKSGRSAGKAYVYLINPGKAFRSKY 136
 QY 123 SLANYLKNGETSLKPEDFDFTLSKRGIKSRKYDCSMAALTSLHONOSNNNSNNMLRTS 182
 DB 137 ELIAAFBKGDITSLDNDPDTLV-TGRGSPSRREQ-----KPPKKPSPKAPGTGRGRG 189
 QY 183 KKKQDVFPSSSSSELQSRGSLNFTSTLLAKEBGVDDVDFRKYRXPXK-----VT 236
 DB 190 R-----PKSGTGRPRPAAASBEQVYQRVLEKSPGKLVMPFOASPGGGBGGGATT 241
 QY 237 ILKGIPIKTKKGGKSCSGFVQSDSKR-----SVCNKADESEPVNAQKSDRT 287

Db 242 SAQVWIK--RPRKRAKADPPQAI PKRGKPGSVAAAAABAKKAVKESIRSVHET 299
OY 288 VCISDAGACGETLSTVSTSE-----NSLVKKERSLSGSGNFCSBOKTSGIINKFCGAK 340
Db 300 VLPKKRRTRETVSIEVKEVVKPLVSTLGEKSGKGLTKCKSPGRKSKSSPKGRSSSAS 359
OY 341 DSEHNEKYEDTFLSESEBIGTKVEVERKEHLHTDILK-----RGSEMDNNCSP 388
Db 360 SPKKEHH-----HHHSESTKAPMPLPSPPPPEPESSEDPISP 400
OY 389 TRKDFTEKIFQEDTIPR 406
Db 401 PEPDLSSSICKKEMPR 418

Search completed: August 22, 2005, 10:09:03
Job time : 182 secs

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OM protein - protein search, using sw model

Run on: August 22, 2005, 10:02:08 ; Search time 42 Seconds
(without alignments)
1328.708 Million cell updates/sec

Title: US-10-629-951-2

Perfect score: 3055
Sequence: 1 MGTGLESLSLGDRCAPTV.....HKAKYHDMWLNENKSLSL 580

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: p1r1:*
2: p1r2:*
3: p1r3:*
4: p1r4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	209	6.8	476	2	S57963 methyl Cpg binding
2	201	6.6	492	2	S41907 methyl-Cpg-binding
3	145	4.7	1301	2	S51323 SAG3 protein - yea
4	145	4.7	3418	1	G02334 breast cancer tumor
5	141.5	4.6	782	2	S27833 thoptry-associated
6	138.5	4.5	500	2	S55785 nucleolar protein
7	137	4.5	560	2	S53382 protein YKR029c ho
8	136.5	4.5	598	2	B40713 cyclin I - human
9	136.5	4.5	1359	2	T34036 hypothetical prote
10	135	4.4	822	2	T41622 probable ABC trans
11	132.5	4.3	286	2	A72882 probable A/G-spect
12	132	4.3	1819	2	A71928 cag island protein
13	131.5	4.3	997	2	T43523 cut1 protein - fi
14	130	4.3	1927	2	G64585 cag pathogenicity
15	128.5	4.2	946	2	A96748 hypothetical prote
16	128	4.2	853	2	T51505 hypothetical prote
17	126	4.1	561	2	H86442 unknown protein li
18	124.5	4.1	491	2	T50346 hypothetical prote
19	124	4.1	650	2	T33350 hypothetical prote
20	124	4.1	1702	2	T14050 protein kinase (EC
21	123	4.0	697	2	C97120 topoisomerase I (l
22	122.5	4.0	669	2	S55024 nebulin, skeletal
23	122	4.0	651	2	C86333 hypothetical prote
24	122	4.0	3924	2	S37431 ankyrin 2, neuroma
25	120.5	3.9	911	2	S51441 hypothetical prote
26	120.5	3.9	1147	2	JN0599 DNA-binding protei
27	120	3.9	1233	2	S56371 hypothetical prote
28	120	3.9	1131	2	A49393 activator 1 large
29	120	3.9	1440	2	T33813 hypothetical prote

30	120	3.9	1658	2	S55101 hypothetical prote
31	120	3.9	2253	2	T30336 nuclear/mitotic ap
32	119.5	3.9	1740	2	T43773 hypothetical prote
33	119.5	3.9	3122	2	T17202 DNA-directed DNA p
34	119	3.9	646	2	F71620 hypothetical prote
35	119	3.9	797	2	H04919 hypothetical prote
36	119	3.9	991	2	H06168 hypothetical prote
37	119	3.9	1002	2	T30546 major surface glyc
38	118.5	3.9	891	2	B84614 hypothetical prote
39	118.5	3.9	1040	2	E71412 hypothetical prote
40	118	3.9	533	2	B84590 hypothetical prote
41	118	3.9	734	2	B42680 nucleolar-cytoplas
42	118	3.9	1200	2	A46194 neurofilament prot
43	118	3.9	1377	2	T51447 transcription regu
44	118	3.9	3329	2	T42205 breast cancer susc
45	118	3.9	3329	2	T30904 breast cancer tumor

ALIGNMENTS

RESULT 1

S57963 methyl Cpg binding protein 2 - human (fragment)
C:Species: Homo sapiens (man)
C>Date: 19-Mar-1997 #sequence_revision 25-Apr-1997 #text_change 05-Nov-1999
C:Accession: S57963
R: d'Esposito, M.; Quaderi, N.A.; Ciccocioppa, A.; Bruni, P.; Esposito, T.; D'Urso, M.; B
submitted to the EMBL Data Library, July 1995
A:Description: Physical mapping and expression analysis of an X-linked gene encoding a m
A:Reference number: S57963
A:Accession: S57963
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-476 <DE>
A:Cross-references: EMBL:X89430; NID:G899295; PIDN:CAA61599.1; PID:G899296

Query Match 6.8% Score 209; DB 2; Length 476;
Best Local Similarity 24.0% Pred. No. 6.5e-06;
Matches 104; Conservative 53; Mismatches 149; Indels 128; Gaps 17;

QY	37	EDVAMELEBRVGEDEBQNMIKRSSECNPLLOEBIASAOPG-----ATAGTE	81
DB	13	KDXPLKFKYKKDKKEKEGHEPVQSAHSAEBAKAKTSESGSARLCEASASP	72
QY	82	CRKSV-----PCGERVVKQPLRGKTNGRPDVFIFISQGLKFKRKSILANTL	128
DB	73	QRRSIIIRDGPWYDDPTLPFGWTRKLKQKSGRSAGKYDVYLINQKAFRSKVELIAYF	132
QY	129	HONGETSLKPEPFDFTVLSKGIKRSRYDCSMAALTSHLQNSNNNNLFRSKCKKDV	188
DB	133	EKVGPDTSLDPNDFDTV-TGRGSPSR-----RKQKPPK--	165
QY	189	FMPSSSSBLQESRGL--SNFTSHLLKDEGDVDVFRKVRKPKGVTLIKGIPRK	245
DB	166	---PKSPAPRGGRGRGKRGKSGGTRPKAATSEGVQK--RLLEKSPK--LLVMQPF-Q	217
QY	246	TKKGRKSCSGFVQSDS---KRESVCKADAESPPVAOKSOLDRTVCISDAGAGETIS	301
DB	218	TPPGKAEGGAGATTSTQWVVIKRPGRKRAEADPOAIPKR-----GRKGSVVA	267
QY	302	VTSSENSLVKKERLSGSSNFCSEQKTSGLINKCSAKDSHNEKYEPTFLESEIGTK	361
DB	266	AAAEAKKKAIVESIR-----SVQETVLPFKK-----RTMET-----VS	303
QY	362	VEVERKEHLHTDIL--KRGSEMDNNSPTRKDFTEKIFQEDTTPRQIERRKTSLYPS	419
DB	304	IEVKEVVKPLVSTLGEKSGKGLKTKCKSPGR-----SKESSPKGR-----	344
QY	420	SKYNKALSPPRK 433	
DB	345	---SSASASPKE 355	

N/Alternate names: breast cancer susceptibility protein BRCA2
 C/Species: Homo sapiens (man)
 C/Date: 21-Dec-1996 #sequence_revision 06-Jun-1997 #text_change 09-Jul-2004
 C/Accession: G02334; S68501
 R/Taxid: 9606; R/Accession: G02334; S68501
 R/Author: S.V.; R/Comment: J.M.; Couch, F.J.; Neuhausen, S.; Bell, R.; Berry, S.; Bogde, M.; Snyder, S.; Stetefeld, M.; Stroup, C.; Swedlund, B.; Teng, D.; Thomas, A.; Truett, M.; et al. The EMBL Data Library, December 1995
 A/Reference number: H01078
 A/Accession: G02334
 A/Status: translated from GB/EMBL/DBJ
 A/Molecule type: mRNA
 A/Residues: 1-3418 <TAN>
 A/Cross-references: UNIPROT:P51587; EMBL:U43746; NID:G1161383; PDB:1ABO7223.1; PID:G1161383
 R/Author: R.; Bignelli, G.; Lancaster, J.; Swift, S.; Seal, S.; Mangion, J.; Collins, N.; Smith, A.; Connor, F.; Arason, A.; Gudmundsson, J.; Plenc, D.; Keisell, D.; Ford, D.; To, N.; et al. Nature 378, 789-792, 1995
 A/Author: Futreal, P.A.; Ashworth, A.; Stratton, M.R.
 A/Title: Identification of the breast cancer susceptibility gene BRCA2.
 A/Reference number: S68501; MUID:96112016; PMID:8524414
 A/Accession: S68501
 A/Molecule type: mRNA
 A/Residues: 282-371, 'N', 373-598, 'S', 600-1108, 'EQ', 1111-1119, 'D', 1121-2321, 'V', 2323-2386, 'C', 'Genetics':
 A/Genes: GDB:BRCA2
 A/Cross-references: GDB:387848; OMIM:600185
 A/Map position: 13q12.3-13q12.3
 C/Superfamily: DNA recombination repair protein, BRCA2 type
 C/Keywords: polymorphism; tumor suppressor

Query Match 4.7%; Score 145; DB 1; Length 3418;
 Best Local Similarity 20.9%; Pred. No. 1.2;
 Matches 96; Conservative 67; Mismatches 158; Indels 138; Gaps 23;

```

QY 36 KEDVAMELRVGEDEQMMIKSSSECNPLQEPISAOGCATGTGTCRKSVPQGWYVK 95
DB 972 KSDISLNDIKIPKKNNDYNNKAWG-----LGGISNHSFGSGSRRTASNEIKLSENNIK 1026
QY 96 ORLFGTAGRFVDYFISPOGLKFRSKSLANYLHKAGETSL--KPEDDFTVLSKRGIKSR 154
DB 1027 SKGFPR-----DLEBQYPTSL--ACVEIVNTLALNOKKLSKPSIN--TVSAHLQSSV 1077
QY 155 YKDCSMAALITSLH--ONQSNNSNMNLTMTSKCKKQVFMPPSSSELOESRGLSNFTSTH 212
DB 1078 VSDCKSHITPQWLFQKQPDNSNHL-----TPSQKALITE-----LSTTL 1118
QY 213 LKEDGCVDDVNRKTRK-----PKGKVTILK----- 239
DB 1119 -----EESGSGFETPTQFKPSYILQKSTFEVENQMTLKTSTSECRDADLHVIMAPSIG 1174
QY 240 -----GPIPK-----TKGCRKSCSGFVQSDSKRESVCNKADAESEFPAQKSQL 284
DB 1175 QVDSKQFQGTGTAIKKFKAGLKNCKKASGLITB-----NEVGPRGFSAGTKL 1227
QY 285 D-RTVCISDAGAGETLSTVSEENSLVKKKERSLSSGNSFCSEQKTSGLIINKECSAKDS 343
DB 1228 NVSTALQKAVKLFSDIENISETS--AEVHPISLSSSKCHD-----SVVSMF--KLEN 1277
QY 344 HNEKEDTFLSESE-----IGKRVKVERKEHLHDLKRGSEMMN--CSP 388
DB 1278 HNRK-----TVSEKNNKCOLIKNNIEMTTGTVEEI-----TENKRNTEENDKYTYA 1327
QY 389 TRK-----DFTGEKTFQEDTIPRQIERRKSLYFSSKYN 423
DB 1328 SRNSHNLFFQSDSSSKNDIV---CIHQEDTDLFTQDN 1363

```

RESULT 5
 S27833
 rhoptry-associated protein 1 precursor - malaria parasite (Plasmodium falciparum)
 N/Alternate names: protective antigen
 C/Species: Plasmodium falciparum
 C/Date: 17-Apr-1993 #sequence_revision 17-Apr-1993 #text_change 09-Jul-2004

C/Accession: A45514; S27833
 R/Ridley, R.G.; Takacs, B.; Lahm, H.W.; Delves, C.J.; Goman, M.; Certa, U.; Matile, H.; Mol. Biochem. Parasitol. 41, 125-134, 1990
 A/Title: Characterisation and sequence of a protective rhoptry antigen from Plasmodium falciparum
 A/Reference number: A45514; MUID:90348711; PMID:2200961
 A/Accession: A45514
 A/Status: preliminary
 A/Molecule type: DNA
 A/Residues: 1-782 <RT>
 A/Cross-references: UNIPROT:Q26007; GB:M32853; NID:G160656; PID:G160657
 C/Superfamily: Plasmodium falciparum rhoptry-associated protein 1

Query Match 4.6%; Score 141.5; DB 2; Length 782;
 Best Local Similarity 21.1%; Pred. No. 0.27;
 Matches 110; Conservative 71; Mismatches 176; Indels 165; Gaps 25;

```

QY 123 SLANTYLNKGER-----SLKPEDFDE-----TVLSKRGIKRKYKDCSMAALITSLONQS 171
DB 18 NVADGGINNGNNYKTIINDFNPDVNYMTPIKKEFLNSYED--EFSSESFLNKS 75
QY 172 -NNSNMNLTMTSKCKQVFMPPSSSELOESRGLSNFTSTHLLKEDGCVDDVNRKTRK 230
DB 76 VDDGININDTSTSNK-----SSKKHGSRVRSASAAALTEEDSKODMEF----- 122
QY 221 PKGKVTILKGIPIKTKKGRKSCSGFVQSDSKRESVCNKADAESEFPAQKSQLDRTVCI 290
DB 123 -KASPSVKTSTPSGTOGLKSS--PSTKSSPSN-----V 158
QY 291 SDGACGETLSTVSEENSLVKKERSLSSGNSFCSEQKTSGLIINKECSAKDSHNEKYED 350
DB 159 KSASPHGS-----NSSEESTTKSKRSAS-----VAGIV-----GADBEAPPAPKN 200
QY 351 TPLESEBI-GTKVVERKEHLHT-----DILKRS-----EMDNCSPTKQDFG 395
DB 201 TLTPLRELYPTVNLFNKYSLNMEENINILKNQDGLVAQKEPEYDENNEKAQODKK 260
QY 396 --EKI-----FOEDTIPRQIERRKSLYFSSKYNKALSPPRKAFKMTTPRS 443
DB 261 ALEKIGKQSDPEPFSEKFLNQKENVAGSRSRFSK--LNP-----PKK----- 307
QY 444 PNLVQETLFHPWLLIATFLNRTSGMALPVLKLEKYPKA----- 488
DB 308 -DEVIEKT-----EVSCKTSGIGFNLTDKAKYLGATGYQVEPTMLYNCNNNLFDIT 363
QY 489 -----EVAKTDWRDVEBLKPLGLYDLRAK--TVKSDSEITLQMKYPIEL 534
DB 364 ESLQGRITIDIKKRESMISTTFEQKCEKLNMGVLLLEINDTQCKEFT----- 410
QY 535 HGIGKYGNDYRIF--CVNEMKQVHP-----EDHKLNKYH 567
DB 411 -CIGSGFGEHRLRYFENDLFRPHNIDYLLTLADYKQLQKNH 451

```

RESULT 6
 S55785
 nucleolar protein gar2 - fission yeast (Schizosaccharomyces pombe)
 N/Alternate names: probable RNA-binding protein RBD18
 C/Species: Schizosaccharomyces pombe
 C/Date: 28-Oct-1996 #sequence_revision 07-Feb-1997 #text_change 16-Aug-2004
 C/Accession: S55785; S68087; T37634; S52338
 R/Gullit, M.P.; Girard, J.P.; Zabetakis, D.; Lapeyre, B.; Melese, T.; Calergues-Ferrer, Nucleic Acids Res. 23, 1912-1918, 1995
 A/Title: gar2 is a nucleolar protein from Schizosaccharomyces pombe required for 18S rRNA biogenesis
 A/Reference number: S55785; MUID:95319932; PMID:7596817
 A/Accession: S55785
 A/Status: nucleic acid sequence not shown
 A/Molecule type: DNA
 A/Residues: 1-500 <GUL>
 A/Cross-references: UNIPROT:P41891; EMBL:Z48166; NID:G663261; PDB:1CAA8179.1; PID:G663261
 Curr. Genet. 29, 307-315, 1996
 A/Title: Molecular analysis of a novel Schizosaccharomyces pombe gene containing two RNUA1-like domains
 A/Reference number: S68087; MUID:96171513; PMID:8598051

A:Accession: S68087
A:Status: nucleic acid sequence not shown; not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 374-407 <VAN>
R:Brown, D.; Churcher, C.M.; Barrell, B.G.; Rajandream, M.A.; Wood, V.
submitted to the EMBL Data Library, September 1997
A:Reference number: 221733
A:Accession: 137634
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-338, 'S', 340-500 <ERO>
A:Cross-references: EMBL:Z99091; PDB:CA811772.1; GSPDB:GN00066; SPDB:SPAC13F5.09
A:Experimental source: strain 972h-; cosmid G13F5
C:Genetics:
A:Gene: gar2; SPDB:SPAC13F5.09
A:Map position: 1
C:Superfamily: ribonucleoprotein repeat homology
F:264-331/Domain: ribonucleoprotein repeat homology <RM1>
F:367-433/Domain: ribonucleoprotein repeat homology <RM2>

Query Match 4.5%; Score 138.5; DB 2; Length 500;
Best Local Similarity 23.7%; Pred. No. 0.23;
Matches 74; Conservative 50; Mismatches 147; Indels 41; Gaps 11;
Qy 185 KQDVMPSSSSSEIQSRGLSNFTSTHLLKEDGVDPVNFRRKPKKVTTLKGIPIK 244
Db 19 KKGATKEPSKSKITKEAKE-----IAQSKTDVSPKSKKAKRASPE--PSK 68
Qy 245 KTKGCRKSCSGFVQSDSKRESVCNKADAEPPVAQKQDLRTVCTISAGACGELSTTS 304
Db 69 KSVKKQKSKK---KESSSSSESSSSSESSSSSESSSSSS--EBSSESSSESSSS 122
Qy 305 EENSIVK---KKERLSGSGNFCEQKTSGLINKCSAKDSEHNKCYDTPLESEIGTK 361
Db 123 EEVAVTKBEKKSSSESSSSSESEEEEAAY-KIEKKESSSDSSSSSESSSSSS 181
Qy 362 VEEVERKEHLH-TDILKRGSEMDNCSPTRKDFTEKIFQEDTIPRTQIERKTSLYPSS 420
Db 182 SESEEEVEVEKTEKKESSESSSDSSSE-----SGSDSSSESSSS--SE 234
Qy 421 KYNKALSPRRKAPKMTTPRSPFNLYOETLFHDPMLLATIFLANTSGMALPVLWK 480
Db 235 DEKRRKAPASEERPAKITPQSDSN---ET-----CTYFVGLSNWVDDQWLQ 281
Qy 481 FLEKTPSAEVAR 492
Db 282 EPEEYGTIVGAR 293

RESULT 7

S53382
protein YKR029c homolog YUL105w - Yeast (Saccharomyces cerevisiae)
N:Alternate names: hypothetical protein J0819
C:Species: Saccharomyces cerevisiae
C:Date: 05-May-1995 #sequence_revision 01-Sep-1995 #text_change 09-Jul-2004
C:Accession: S53382; S56883; S57163
R:Raamussen, S.W.
submitted to the EMBL Data Library, February 1995
A:Description: A 37.5 kb region of yeast chromosome X includes the SME1, ME2, GSH1 and
A:Reference number: S53376
A:Accession: S53382
A:Molecule type: DNA
A:Residues: 1-560 <RAS>
R:Raamussen, S.W.
submitted to the Protein Sequence Database, September 1995
A:Reference number: S56876
A:Accession: S56883
A:Molecule type: DNA
A:Residues: 1-560 <RAW>
A:Cross-references: EMBL:Z49380; NID:g1008285; PID:g1008286; MIPS:YUL105w
R:Raamussen, S.W.
Yeast_11, 873-883, 1995

A:title: A 37.5 kb region of yeast chromosome X includes the SME1, ME2, GSH1 and CSD3 ge
A:Reference number: S57357; MUID:96090136; PMID:7483851
A:Accession: S57363
A:Status: nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-560 <RAP>
A:Cross-references: EMBL:X85021; NID:g728698; PDB:CAA59389.1; PID:g728705
A>Note: the nucleotide sequence was submitted to the EMBL Data Library, February 1995
C:Genetics:
A:Cross-references: SGD:S0003641
A:Map position: 10U

Query Match 4.5%; Score 137; DB 2; Length 560;
Best Local Similarity 18.7%; Pred. No. 0.33;
Matches 90; Conservative 80; Mismatches 161; Indels 150; Gaps 24;
Qy 118 FRKSSILANTLHKNGETSLKPEPDFVYLKRGTKSKYKCSMAALTSHL--QKQSNNSN 175
Db 82 FQNDRGIFNHSSSGSSKT-----ASTNKRGIAAVALATATPFPPLKKQNDNSK 134
Qy 176 WNLTRSKCKDVMPSSSSSEIQSRGLSNFTSTHLLKEDGVDPVNFRRKPKKGV 235
Db 135 VSV-TNNESSKENKITTSMAE-----DNKPNNGCICSSDSKDEL----- 174
Qy 236 TILKGIPIKTKGCRKSCGFVQSDS-KRESVCNKADAESEPPVAQKQDL----- 285
Db 175 ----FICNCKCTWQHLCYAFKKS DPIKDPVCKRCDSPTK--VQNVQVPMI FPRMG 228
Qy 286 -----RTYCISPAKCGE-----TLSTSENSLVK-KESLS 318
Db 229 DERLFQSSIVTTSASNTNHOQSNNIEOPKKRQLHYAPTTENSNSIRKRLRQKLV 288
Qy 319 SGNFCEQKTSGLINKCSAKD-----SEHNKCYEDTPLESEIGTKVEVERKEH 370
Db 289 VSHFLEK-----ILNVSSNDTEFRAITISEYKDYKVMKFDN-----H 329
Qy 371 LHTDIL-----KRGEMDNCSPTRKDFTEKIFQEDTIPRTQ-IERKTSLYPSSKY 422
Db 330 YDDDWVVCNWSSESRADIEVRKSSNERDF--GVFAADSCVKGELIQEYLGKIDFQKNY 386
Qy 423 NKEALSPRRKAPKMTTPRSPFNLYOET---LFHDPMLLATIFLANTSGMALPVL 478
Db 387 Q-----TDPNNDRYLMGTTPKXVLFHPHPLYIDS--REFG-----L 422
Qy 479 WKFLK--VPSAEVARTADRDVSELKPLGLYDLAKTIVK-----SDPYLTKQWKYP 531
Db 423 TRYIRSCBPVELVTV--RPLDE--KPRGNDCKRKYFVLRALRIDRKGEISVENQWD 477
Qy 532 I 532
Db 478 L 478

RESULT 8

B40713
cyclin I - human (fragment)
C:Species: Homo sapiens (man)
C:Date: 12-May-1994 #sequence_revision 12-May-1994 #text_change 09-Jul-2004
C:Accession: B40713; S35920
R:Hees, H.; Heid, H.; Franke, W.W.
J. Cell Biol. 122, 1043-1052, 1993
A:title: Molecular characterization of mammalian cyclin, a basic protein of the sperm h
A:Reference number: A40713; MUID:93359502; PMID:8354692
A:Accession: B40713
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-598 <HES>
A:Cross-references: UNIPROT:P35663; GB:Z22780; NID:g396104; PDB:CAA80457.1; PID:g396105
C:Keywords: cytoskeleton

Query Match 4.5%; Score 136.5; DB 2; Length 598;
Best Local Similarity 23.3%; Pred. No. 0.39;
Matches 111; Conservative 72; Mismatches 205; Indels 89; Gaps 21;

QY 383 ---DNNCSPTRKDFTGEXI FQEDTIPRT 407
: ||| | : |||
Db 752 HSGENCs-----FLPATVPTT 768

Search completed: August 22, 2005, 10:06:00
Job time : 46 secs

GenCore version 5.1.6
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OM protein - protein search, using SW model

Run on: August 22, 2005, 10:02:08 ; Search time 170 Seconds
(without alignments)
1319.535 Million cell updates/sec

Title: US-10-629-951-2

Perfect score: 3055
Sequence: 1 MGTGLESLSIGDRGAPTV.....HKLNKTHMLWENHEKLSLS 580

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 2105692 seqs, 386760381 residues

Total number of hits satisfying chosen parameters: 2105692

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_16Dec04:*

1: geneseqp19808:*
2: geneseqp19908:*
3: geneseqp20008:*
4: geneseqp20018:*
5: geneseqp20028:*
6: geneseqp20038:*
7: geneseqp20038:*
8: geneseqp20048:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	3055	100.0	580	2 AAW74473	Aaw74473 Human MED
2	3055	100.0	580	7 ADD89906	Add89906 Human 5-m
3	3055	100.0	580	8 ADK13954	Adk13954 Human met
4	2958.5	96.8	565	8 AAY44504	Aay44504 Human del
5	2194.5	71.8	439	2 AAY76548	Aay76548 Human ova
6	1639	53.6	307	7 ADD89915	Add89915 Human 5-m
7	1230	40.3	257	2 AAW88701	Aaw88701 Secreted
8	1230	40.3	257	4 ABB50468	Abbs50468 Human sec
9	1230	40.3	257	6 ABO44725	Abos44725 Novel hum
10	1230	40.3	257	7 ABO26205	Abos26205 Human pro
11	1106	36.2	202	7 ADP89916	Adp89916 Human 5-m
12	880.5	28.8	416	7 ADD89908	Add89908 Chicken 5
13	816	26.7	147	7 ADD89917	Add89917 Human 5-m
14	357	11.7	68	5 AAE22568	Aae22568 Human MBD
15	254	8.3	50	4 ABB51147	Abbs51147 Human sec
16	254	8.3	50	6 ABO45404	Abos45404 Novel hum
17	254	8.3	50	7 ABO26884	Abos26884 Protein a
18	220.5	7.2	467	8 ADK13968	Adk13968 Frog meth
19	220.5	7.2	467	8 ADK13967	Adk13967 Frog meth
20	213.5	7.0	486	8 ADK13972	Adk13972 Human met
21	209	6.8	476	8 ADJ68460	Adj68460 Human hea
22	209	6.8	476	8 ADK13973	Adk13973 Human met
23	208.5	6.8	477	8 ADK13961	Adk13961 Human met
24	208.5	6.8	486	8 ADK13966	Adk13966 Human met
25	208.5	6.8	486	8 ADK13971	Adk13971 Human met

26	208.5	6.8	486	8 ADK13957	Adk13957 Human met
27	208.5	6.8	486	8 ADK13960	Adk13960 Human met
28	208.5	6.8	486	8 ADK13953	Adk13953 Human met
29	208.5	6.8	486	8 ADK13963	Adk13963 Human met
30	208.5	6.8	486	8 ADK13970	Adk13970 Human met
31	208.5	6.8	486	8 ADK13958	Adk13958 Human met
32	208.5	6.8	486	8 ABM82465	Abm82465 Tumour-as
33	208.5	6.8	560	6 ABG72561	Abg72561 TAR dwt-M
34	208.5	6.8	561	6 ABG72557	Abg72557 Mecp2-Tat
35	202.5	6.6	345	8 ADK14010	Adk14010 Chicken m
36	201	6.6	492	8 ADK13964	Adk13964 Rat methy
37	198.5	6.5	484	8 ADK13959	Adk13959 Mouse met
38	198.5	6.5	484	8 ADK13956	Adk13956 Human met
39	198.5	6.5	484	8 ADK13969	Adk13969 Mouse met
40	198.5	6.5	484	8 ADK13965	Adk13965 Mouse met
41	194	6.4	219	3 AAG02051	Aag02051 Human sec
42	186	6.1	68	5 AAE22566	Aae22566 Human Mec
43	186	6.1	92	8 ADK13962	Adk13962 Human met
44	159.5	5.2	744	8 ADP49805	Adp49805 Human 193
45	148.5	4.9	326	3 AAG16859	Aag16859 Arabidops

ALIGNMENTS

RESULT 1
AAW74473 standard; protein; 580 AA.
ID AAW74473

AC AAW74473;

DT 19-MAY-1999 (first entry)

DE Human MED1 endonuclease protein sequence.

EN Endonuclease; MED1; human; methyl-CpG binding endonuclease-1;

KW DNA fidelity; DNA manipulation; cancer; fragile X syndrome; therapy;

KW myotonic dystrophy; Huntington's disease; spinocerebellar ataxia;

KW Kennedy's disease; triplet repeat expansion disorder.

OS Homo sapiens.

PN WO9904626-A1.

PD 04-FEB-1999.

PF 28-JUL-1998; 98WO-US015828.

PR 28-JUL-1997; 97US-0053936P.

PA (FOXC-) FOX CHASE CANCER CENT.

PI Bellacosa A;

DR MPI: 1999-142462/12.

DR N-PSDB; AAX22002.

PT New nucleic acid encoding human endonuclease MED1 involved in DNA

PT mismatch repair - used for diagnosing susceptibility to cancer and

PT fragile X syndrome, and therapeutically.

PS Claim 8; Fig 3; 109pp; English.

This sequence is the human MED1 endonuclease of the invention. MED1 (for methyl-CpG binding endonuclease-1) is used to screen for specific CC modulators (potential therapeutic agents particularly mimetics of MED1) and to study interactions involved in maintaining DNA fidelity, for DNA manipulation and to raise antibodies. Susceptibility or predisposition to cancer (particularly colorectal cancer), or its prognosis, where caused by CC alterations in the MED1-encoding gene, are identified by sequence comparison, amplification, detecting altered polypeptide, and restriction fragment mapping, hybridisation (particularly to probes specific for a

CC mutant allele). These same methods can also be used to diagnose fragile X
CC syndrome and other diseases (e.g. myotonic dystrophy, Huntington's
CC disease, spinocerebellar ataxia and Kennedy's disease), associated with
CC triplet repeat expansion. The DNA, or its fragments, are used as probes
CC and primers in the above diagnostic methods, also to isolate homologous
CC sequences, as sources of antisense sequences and for gene transfer,
CC particularly to restore drug sensitivity to drug-resistant cancer cells
XX

Sequence 580 AA;

Query Match 100.0%; Score 3055; DB 2; Length 580;
Best Local Similarity 100.0%; Pred. No. 7,7e-287;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```
QY 1 MGTGLESLSLGDRAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEQMMIKRSSE 60
DB 1 MGTGLESLSLGDRAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEQMMIKRSSE 60
QY 61 CNPLLOEPISAOAGATAGTECRKSVPCGWERVVKORLFGKTAGRFVYFISPOGLKFRS 120
DB 61 CNPLLOEPISAOAGATAGTECRKSVPCGWERVVKORLFGKTAGRFVYFISPOGLKFRS 120
QY 121 KSLIANYLHKNGETSLKPEDFDFTVLSKRGIKSRKYDCSMAALTSHLQNSNNSNMNLT 180
DB 121 KSLIANYLHKNGETSLKPEDFDFTVLSKRGIKSRKYDCSMAALTSHLQNSNNSNMNLT 180
QY 181 RSKCKDVFMPPSSSELQESRGLSNFTSTHLLKEDGVDDVNRKVRKRGKVTILKG 240
DB 181 RSKCKDVFMPPSSSELQESRGLSNFTSTHLLKEDGVDDVNRKVRKRGKVTILKG 240
QY 241 IPIKTKKGRKSCSGFVQSDSKRESVCKADASEPVAOKSOLDRTVCISDAGACGRTL 300
DB 241 IPIKTKKGRKSCSGFVQSDSKRESVCKADASEPVAOKSOLDRTVCISDAGACGRTL 300
QY 301 SVTSEENSLVKKKERSLSSGSNFCSEQKTSGLINKFCSAKDESHNEKEDTFLSESEIGT 360
DB 301 SVTSEENSLVKKKERSLSSGSNFCSEQKTSGLINKFCSAKDESHNEKEDTFLSESEIGT 360
QY 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFGEEKI FQEDTTPRTOIERRKTSLYSS 420
DB 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFGEEKI FQEDTTPRTOIERRKTSLYSS 420
QY 421 KYNKEALSPPRRKAFKMTTPRSPFNLYOETLFHDPATLLATIFLNTSGKMAIPVLMK 480
DB 421 KYNKEALSPPRRKAFKMTTPRSPFNLYOETLFHDPATLLATIFLNTSGKMAIPVLMK 480
QY 481 FLEKTPSAEVARATADWRDVSELKPLGLYDLRAKTIKVFSDBYLTQWKYPIELHGIGKY 540
DB 481 FLEKTPSAEVARATADWRDVSELKPLGLYDLRAKTIKVFSDBYLTQWKYPIELHGIGKY 540
QY 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHMLWENHEKLSLS 580
DB 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHMLWENHEKLSLS 580
```

RESULT 2

ADD89906 ADD89906 standard; protein; 580 AA.

ADD89906;

29-JAN-2004 (first entry)

Human 5-methylcytosine DNA glycoylase.

Human; 5-methylcytosine DNA glycoylase; enzyme; Cpg.

Homo sapiens.

WO2003078593-A2.

25-SEP-2003.

XX

PF 14-MAR-2003; 2003WO-US007933.

XX 15-MAR-2002; 2002US-0364689P.

XX (EPIG-) EPIGENOMICS AG.

PA Lofton-Day CE, Day JK;

PI WPI; 2003-779127/73.

DR N-PSDB; ADD89905.

PT Labeling methylated or methylatable Cpg sequences, useful e.g. for
diagnostic detection of altered methylation, comprises replacing

PT methylated cytosine by labeled cytosine.

XX Claim 11; Page 53-55; 73pp; English.

PS The present sequence is the protein sequence of human 5-methylcytosine
XX DNA glycoylase (5-MCDG). The enzyme acts by cleaving glycosyllic bonds at
XX methylated Cpg sites of DNA, removing 5-methylcytosine from the DNA
XX backbone as a free base. Human 5-MCDG can be used in a claimed method for
XX labelling Cpg sequences corresponding to methylated Cpg sequences in an
XX isolated DNA sample. The method comprises: digesting the genomic DNA with
XX a restriction endonuclease to produce genomic DNA fragments; treating the
XX genomic DNA fragments with 5-MCDG such that one or more 5-methylcytosine
XX bases are removed to produce abasic genomic DNA fragments; and treating
XX these abasic genomic DNA fragments with base excision repair enzymes in
XX the presence of labelled dCTP such that 5-methylcytosine removed from the
XX genomic DNA fragments is replaced by labelled cytosine in the
XX one or more corresponding positions of the abasic genomic DNA fragments
XX to produce labelled genomic DNA fragments, so that specific labelling of
XX Cpg sequences corresponding to methylated Cpg sequences is achieved. The
XX 5-MCDG is also used in a claimed method for comparing Cpg methylation
XX status, extent or pattern between or among reference and test genomic DNA
XX samples, and in a claimed method for labelling potentially-methylatable
XX Cpg sequences in Cpg-containing genomic DNA fragments. The methods are
XX used to identify methylated and/or potentially methylatable Cpg
XX dinucleotides in genomic DNA, including comparison of methylation pattern
XX between healthy and diseased samples, for diagnosis.

Sequence 580 AA;

Query Match 100.0%; Score 3055; DB 7; Length 580;
Best Local Similarity 100.0%; Pred. No. 7,7e-287;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```
QY 1 MGTGLESLSLGDRAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEQMMIKRSSE 60
DB 1 MGTGLESLSLGDRAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEQMMIKRSSE 60
QY 61 CNPLLOEPISAOAGATAGTECRKSVPCGWERVVKORLFGKTAGRFVYFISPOGLKFRS 120
DB 61 CNPLLOEPISAOAGATAGTECRKSVPCGWERVVKORLFGKTAGRFVYFISPOGLKFRS 120
QY 121 KSLIANYLHKNGETSLKPEDFDFTVLSKRGIKSRKYDCSMAALTSHLQNSNNSNMNLT 180
DB 121 KSLIANYLHKNGETSLKPEDFDFTVLSKRGIKSRKYDCSMAALTSHLQNSNNSNMNLT 180
QY 181 RSKCKDVFMPPSSSELQESRGLSNFTSTHLLKEDGVDDVNRKVRKRGKVTILKG 240
DB 181 RSKCKDVFMPPSSSELQESRGLSNFTSTHLLKEDGVDDVNRKVRKRGKVTILKG 240
QY 241 IPIKTKKGRKSCSGFVQSDSKRESVCKADASEPVAOKSOLDRTVCISDAGACGRTL 300
DB 241 IPIKTKKGRKSCSGFVQSDSKRESVCKADASEPVAOKSOLDRTVCISDAGACGRTL 300
QY 301 SVTSEENSLVKKKERSLSSGSNFCSEQKTSGLINKFCSAKDESHNEKEDTFLSESEIGT 360
DB 301 SVTSEENSLVKKKERSLSSGSNFCSEQKTSGLINKFCSAKDESHNEKEDTFLSESEIGT 360
QY 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFGEEKI FQEDTTPRTOIERRKTSLYSS 420
DB 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFGEEKI FQEDTTPRTOIERRKTSLYSS 420
```

QY 421 KYNKALSPRRKAFKKTTPRRSPNNLVOETLFHDPWKLLIATITFLNRTSGMAIPVLMK 480
 DB 421 KYNKALSPRRKAFKKTTPRRSPNNLVOETLFHDPWKLLIATITFLNRTSGMAIPVLMK 480
 QY 481 FLEKTPSAEAVARTADWRDVSLLKPLGLYDLRAKTIIVKPSDEYLTQWKYPIELHGIGKY 540
 DB 481 FLEKTPSAEAVARTADWRDVSLLKPLGLYDLRAKTIIVKPSDEYLTQWKYPIELHGIGKY 540
 QY 541 GNDSTRICVNMKOVHPEDHKLANKYHDMLENHEKLSLS 580
 DB 541 GNDSTRICVNMKOVHPEDHKLANKYHDMLENHEKLSLS 580

RESULT 3

ADK13954 standard; protein; 580 AA.

ADK13954

ADK13954; (first entry)

03-JUN-2004 (first entry)

Human methyl-CpG-binding protein #9.

Ret syndrome; methyl-CpG-binding protein 2; MECP2; neurodevelopmental disease; autism; non-syndromic mental retardation; idiopathic neonatal encephalopathy; idiopathic infantile spasms; idiopathic cerebral palsy; Angelman syndrome; schizophrenia; human.

Homo sapiens.

US6709817-B1.

23-MAR-2004.

07-SEP-2000; 2000US-00657013.

07-SEP-1999; 99US-0152778P.

(BAYU) BAYLOR COLLEGE MEDICINE.

Zoghbi HY, Van Den Veyver IB, Amir R, Francke U;

MPI; 2004-256068/24.

Screening human for Rett syndrome comprises detecting mutation in nucleic acid sequence encoding methyl-CpG-binding protein 2 (MECP2).

Disclosure; SEQ ID NO 56; 125pp; English.

The invention relates to a method of screening a human for Rett syndrome comprising detecting a mutation in a nucleic acid sequence encoding methyl-CpG-binding protein 2 (MECP2). The method is useful for screening a human for Rett syndrome. The method is useful for screening neurodevelopmental diseases such as Rett syndrome, autism, non-syndromic mental retardation, idiopathic neonatal encephalopathy, idiopathic infantile spasms, idiopathic cerebral palsy, Angelman syndrome and schizophrenia. The present sequence represents the amino acid sequence of a methyl-CpG-binding protein.

Sequence 580 AA;

Query Match 100.0%; Score 3055; DB 8; Length 580;

Best Local Similarity 100.0%; Pred. No. 7.7e-287;

Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 MGTTGLESLSLGRGAAPVTSSERLVPDPNDLRKEDVAMELEVRGEDEEQMIRKSS 60

1 MGTTGLESLSLGRGAAPVTSSERLVPDPNDLRKEDVAMELEVRGEDEEQMIRKSS 60

61 CNPLLOEPINSAOGATAGTECKSVPCGWERVVKQMLFKKTAGRPVYFISQGLKFS 120

61 CNPLLOEPINSAOGATAGTECKSVPCGWERVVKQMLFKKTAGRPVYFISQGLKFS 120

61 CNPLLOEPINSAOGATAGTECKSVPCGWERVVKQMLFKKTAGRPVYFISQGLKFS 120

61 CNPLLOEPINSAOGATAGTECKSVPCGWERVVKQMLFKKTAGRPVYFISQGLKFS 120

61 CNPLLOEPINSAOGATAGTECKSVPCGWERVVKQMLFKKTAGRPVYFISQGLKFS 120

61 CNPLLOEPINSAOGATAGTECKSVPCGWERVVKQMLFKKTAGRPVYFISQGLKFS 120

61 CNPLLOEPINSAOGATAGTECKSVPCGWERVVKQMLFKKTAGRPVYFISQGLKFS 120

61 CNPLLOEPINSAOGATAGTECKSVPCGWERVVKQMLFKKTAGRPVYFISQGLKFS 120

61 CNPLLOEPINSAOGATAGTECKSVPCGWERVVKQMLFKKTAGRPVYFISQGLKFS 120

QY 121 KSLANLYHKNGETSLKPEDFTVYLSKRGKSKRYKDCSMAALTSHLQNSNNSMNLRT 180
 DB 121 KSLANLYHKNGETSLKPEDFTVYLSKRGKSKRYKDCSMAALTSHLQNSNNSMNLRT 180
 QY 181 RSKCKGVFMPSSSSSELQESRGLSNFTSTHLLKEDGVDDVNFRRKVRKPKGVYTIK 240
 DB 181 RSKCKGVFMPSSSSSELQESRGLSNFTSTHLLKEDGVDDVNFRRKVRKPKGVYTIK 240
 QY 241 IPIKTKKCKGKSGGFTVQSDSKRESVCKADASEPVAQNSOLDRTVCISDAGCGE 300
 DB 241 IPIKTKKCKGKSGGFTVQSDSKRESVCKADASEPVAQNSOLDRTVCISDAGCGE 300
 QY 301 SVTSENLSLVKKKESLSSGNSPCSEOKTSGIINFCSAKSEHNEKTEDTLESEIGT 360
 DB 301 SVTSENLSLVKKKESLSSGNSPCSEOKTSGIINFCSAKSEHNEKTEDTLESEIGT 360
 QY 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFTEKIFQEDTIPRTQIERKSTLYFS 420
 DB 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFTEKIFQEDTIPRTQIERKSTLYFS 420
 QY 421 KYNKALSPRRKAFKKTTPRRSPNNLVOETLFHDPWKLLIATITFLNRTSGMAIPVLMK 480
 DB 421 KYNKALSPRRKAFKKTTPRRSPNNLVOETLFHDPWKLLIATITFLNRTSGMAIPVLMK 480
 QY 481 FLEKTPSAEAVARTADWRDVSLLKPLGLYDLRAKTIIVKPSDEYLTQWKYPIELHGIGKY 540
 DB 481 FLEKTPSAEAVARTADWRDVSLLKPLGLYDLRAKTIIVKPSDEYLTQWKYPIELHGIGKY 540
 QY 541 GNDSTRICVNMKOVHPEDHKLANKYHDMLENHEKLSLS 580
 DB 541 GNDSTRICVNMKOVHPEDHKLANKYHDMLENHEKLSLS 580

RESULT 4

AA44504 standard; protein; 565 AA.

AA44504

AA44504; (first entry)

27-MAR-2000 (first entry)

Human delta228-UV damage endonuclease.

Delta228-UVDE; ultraviolet damage endonuclease; GST signal peptide; glutathione-S-transferase signal peptide; uvei+ gene product; UV irradiation; DNA damage; UV radiation damage; photoproduct; abasic site; aplatinum diaduct; mismatched nucleotide pairing; nucleotide alkylation; skin cancer.

Homo sapiens.

MO9963828-A1.

16-DEC-1999.

08-JUN-1999; 99WO-US012910.

08-JUN-1998; 98US-0086521P.

18-MAY-1999; 99US-0134752P.

(UYEM-) UNIV EMORY.

Doetsch PW, Kaur B, Avery AM;

MPI; 2000-116417/10.

A new truncated ultraviolet damage endonuclease for treatment of skin cancers.

Claim 16; Page 60; 133pp; English.

The present sequence is human delta228-UV damage endonuclease. Delta228-

The present sequence is human delta228-UV damage endonuclease. Delta228-

The present sequence is human delta228-UV damage endonuclease. Delta228-

The present sequence is human delta228-UV damage endonuclease. Delta228-

The present sequence is human delta228-UV damage endonuclease. Delta228-

CC UVDE is a 288 amino acid deletion of the N-terminal of the uvel+ gene
CC product. This is expressed in frame with a GST leader sequence to
CC generate a fusion protein. This provides stable endonuclease fragments
CC for cleaving a double-stranded DNA molecule that has a distorted
CC structure resulting from UV radiation damage, a photoproduct, an abasic
CC site, mismatched nucleotide pairing, apatidium diaduct, an intercalated
CC molecule or alkylation of a nucleotide. Uvelp can be used in compositions
CC for internal or topical application and as a therapeutic agent for skin
CC cancers

XX Sequence 565 AA;

Query Match 96.8%; Score 2958.5; DB 3; Length 565;
Best Local Similarity 97.4%; Pred. No. 1.7e-277;
Matches 565; Conservative 0; Mismatches 0; Indels 15; Gaps 1;

QY 1 MGTGLSELSLQDRGAFTVTSSERLVDPDPNDLKEDEVAMELEVRGDEEQMMIKRSSE 60
DB 1 MGTGLSELSLQDRGAFTVTSSERLVDPDPNDLKEDEVAMELEVRGDEEQMMIKRSSE 60
QY 61 CNPLLOEPIASAOFATATGTECRKSVPCGWERVYKQRLFGKTAGRFVYFISPOGLKERS 120
DB 61 CNPLLOEPIASAOFATATGTECRKSVPCGWERVYKQRLFGKTAGRFVYFISPOGLKERS 120
QY 121 KSLIANYLHKAGETSLKPEDEFTVLSKRGISKRYKDCSMALTSHLQNSNNSMNLRT 180
DB 121 KSLIANYLHKAGETSLKPEDEFTVLSKRGISKRYKDCSMALTSHLQNSNNSMNLRT 180
QY 121 KSLIANYLHKAGETSLKPEDEFTVLSKRGISKRYKDCSMALTSHLQNSNNSMNLRT 180
DB 121 KSLIANYLHKAGETSLKPEDEFTVLSKRGISKRYKDCSMALTSHLQNSNNSMNLRT 180
QY 181 RSKCKQVFMPPSSSELQESRGLSNFTSTHLLKEDGVDVDFRKYRKPKGKVTIILKG 240
DB 181 RSKCKQVFMPPSSSELQESRGLSNFTSTHLLKEDGVDVDFRKYRKPKGKVTIILKG 240
QY 181 RSKCKQVFMPPSSSELQESRGLSNFTSTHLLKEDGVDVDFRKYRKPKGKVTIILKG 240
DB 181 RSKCKQVFMPPSSSELQESRGLSNFTSTHLLKEDGVDVDFRKYRKPKGKVTIILKG 240
QY 241 IPIKTKKGCRCSCGFVOSDSKRSYCNKADAESEPAOKSOLDRTVCIDAGACGTL 300
DB 241 IPIKTKKGCRCSCGFVOSDSKRSYCNKADAESEPAOKSOLDRTVCIDAGACGTL 300
QY 241 IPIKTKKGCRCSCGFVOSDSKRSYCNKADAESEPAOKSOLDRTVCIDAGACGTL 300
DB 241 IPIKTKKGCRCSCGFVOSDSKRSYCNKADAESEPAOKSOLDRTVCIDAGACGTL 300
QY 301 SVTSENSLVKKKERSLSGSGNFCSEOKTSGIINFCGAKSEHNEKTEDFTLSESEIGT 360
DB 301 SVTSENSLVKKKERSLSGSGNFCSEOKTSGIINFCGAKSEHNEKTEDFTLSESEIGT 360
QY 289 ---SEENSLVKKKERSLSGSGNFCSEOKTSGIINFCGAKSEHNEKTEDFTLSESEIGT 345
DB 289 ---SEENSLVKKKERSLSGSGNFCSEOKTSGIINFCGAKSEHNEKTEDFTLSESEIGT 345
QY 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFTEGKI FQEDTIPRTQIERRKTSLYPSS 420
DB 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFTEGKI FQEDTIPRTQIERRKTSLYPSS 420
QY 346 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFTEGKI FQEDTIPRTQIERRKTSLYPSS 405
DB 346 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFTEGKI FQEDTIPRTQIERRKTSLYPSS 405
QY 421 KYNKEALSPRRKAFKKWTPRSPFNVLVOETLPHDPWKLIAIIFLNTSKMAIPVLMK 480
DB 406 KYNKEALSPRRKAFKKWTPRSPFNVLVOETLPHDPWKLIAIIFLNTSKMAIPVLMK 465
QY 481 FLEKYPSEAVARTADWRDVSELKPLGLYDLRAKTIYFSDVYLTKQWKYPIELHIGIKY 540
DB 466 FLEKYPSEAVARTADWRDVSELKPLGLYDLRAKTIYFSDVYLTKQWKYPIELHIGIKY 525
QY 541 GNDSYRIFCVNEKQVHEDHKLKYNKHDMLWENHEKLSLS 580
DB 526 GNDSYRIFCVNEKQVHEDHKLKYNKHDMLWENHEKLSLS 565

RESULT 5
AAV76548
ID AAV76548 standard; protein; 439 AA.

XX AAV76548;

XX 10-APR-2000 (first entry)

XX Human ovarian tumor EST fragment encoded protein 44.

XX Expressed sequence tag: EST; human; ovarian tumor; anticancer;

KM gene therapy; treatment.

XX Homo sapiens.

XX DE19817557-A1.

XX 21-OCT-1999.
PD 98DE-01017557.
XX 09-APR-1998;
PF 98DE-01017557.
XX 09-APR-1998;
PR 98DE-01017557.
XX (META-) METAGEN GES GENOMFORSCHUNG MBH.
PA
XX Rosenthal A, Specht T, Hinzmann B, Schmitt A, Pilarsky C, Dahl E;
PI WPI; 1999-591920/51.
XX N-PSDB; AA277469.
DR
XX New nucleic acid sequences expressed in ovarian, and some other, cancer
PT tissues, and derived polypeptides, for treatment of ovarian cancer and
PT identification of therapeutic agents.

PS Claim 25; Page 261, 310pp; German.

CC This invention describes novel nucleic acid (cDNA) sequences (A) which
CC have anticancer activity and are highly expressed in ovarian tumor tissue
CC (and some also in testis and breast cancer tissue). The products of the
CC invention can be used for gene therapy. (A) are used (i) for recombinant
CC expression of polypeptides (B) and (ii) to isolate complete genes. (B)
CC are used (i) to identify agents suitable for treatment of ovarian cancer;
CC (ii) directly for treating this form of cancer (including expression from
CC gene therapy vectors) and (iii) for generation of specific antibodies.
CC (A) are identified by assembling ESTs (expressed sequence tags) from a
CC particular tissue type before comparison of expression patterns. This
CC allows a significantly longer fragment of the gene to be revealed, so
CC should reduce the number of failures associated with the fact that ESTs
CC from different libraries may represent different parts of the same
CC unknown gene; distorting the estimated frequency of occurrence in a
CC particular tissue. AAV76505-776638 represent protein fragments encoded by
CC the human ovarian tumor cDNA library derived EST fragments represented in
CC AA277450-277572

XX Sequence 439 AA;

Query Match 71.8%; Score 2194.5; DB 2; Length 439;
Best Local Similarity 96.0%; Pred. No. 1.8e-203;
Matches 427; Conservative 3; Mismatches 6; Indels 9; Gaps 3;

QY 95 KQRLFGTAGRFD-VYRISQGLKFRKSSLIANYLHNGENSLKPEDEFTVLSKRGIKS 153
DB 1 KSLLF--TSKFPPLISFSSPGLKFRKSSLIANYLHNGETSLKPEDEFTVLSKRGIKS 58
QY 154 RYKDCSMALTSHLQNSNNSMNLRTSRKCKKQVFMPPSSSELQESRGLSNFTSTHLL 213
DB 59 RYKDCSMALTSHLQNSNNSMNLRTSRKCKKQVFMPPSSSELQESRGLSNFTSTHLL 118
QY 214 LKEDGVDVDFRKYRKPKGKVTIILKGIPIKTKKGCRCSCGFVOSDSKRSYCNKADA 273
DB 119 LKEDGVDVDFRKYRKPKGKVTIILKGIPIKTKKGCRCSCGFVOSDSKRSYCNKADA 178
QY 274 ESEPVAKQSQUDRVCTSDAGACETLSVTSENSLVKKKERSLSGSGNFCSEOKTSGII 333
DB 179 ESEPVAKQSQUDRVCTSDAGACETLSVTSENSLVKKKERSLSGSGNFCSEOKTSGII 238
QY 334 NKFCSAKDSSEHNEKTEDFTLSESEIGTKVEVERKEHLHTDILKRGSEMDNNSPTRKDF 393
DB 239 NKFCSAKDSSEHNEKTEDFTLSESEIGTKVEVERKEHLHTDILKRGSEMDNNSPTRKDF 298
QY 394 TGEKIFQEDTIPRTQIERRKTSLYFSSSKYNKEALSPRRKAFKKWTPRSPFNVLVOETLF 453
DB 299 T-----EDTIPRTQIERRKTSLYFSSSKYNKEALSPRRKAFKKWTPRSPFNVLVOETLF 352
QY 454 HDPWKLIAIIFLNTSKMAIPVLMKFLKYPSEAVARTADWRDVSELKPLGLYDLRA 513
DB 353 HDPWKLIAIIFLNTSKMAIPVLMKFLKYPSEAVARTADWRDVSELKPLGLYDLRA 412
QY 514 KTIYKFSDEVYLTKQWKYPIELHIGIG 538

DB 413 KTIKFSDEYLTQWKYPLELHGIG 437

RESULT 6
ADD89915
ID ADD89915 standard; protein; 307 AA.
XX
XX ADD89915;
AC
XX 29-JAN-2004 (first entry)
XX
XX Human 5-methylcytosine DNA glycosylase N-terminal deletion mutant.
XX
XX Human; 5-methylcytosine DNA glycosylase; enzyme; Cpg; mutant; mutelin.
XX
XX Homo sapiens.
XX
XX WO2003078593-A2.
XX
XX 25-SEP-2003.
XX
XX 14-MAR-2003; 2003WO-US007933.
XX
XX 15-MAR-2002; 2002US-0364689P.
XX
XX (EPIC-) EPIGENOMICS AG.
XX
XX Lofton-Day CE, Day JK;
XX
XX WPI; 2003-779127/73.
XX
XX Labeling methylated or methylatable Cpg sequences, useful e.g. for
XX PT diagnostic detection of altered methylation, comprises replacing
XX PT methylated cytosine by labeled cytosine.
XX
XX Claim 11; Page 70-71; 73pp; English.
XX
XX The present sequence is the protein sequence of an N-terminal deletion
XX CC mutant of human 5-methylcytosine DNA glycosylase (5-MCDG), in which amino
XX CC acid residue 1 corresponds to amino acid 274 of the full-length protein
XX CC ADD89906. 5-MCDG acts by cleaving glycosylic bonds at methylated Cpg
XX CC sites of DNA, removing 5-methylcytosine from the DNA backbone as a free
XX CC base. The N-terminal deletion mutant shows enhanced deglycosylase
XX CC specificity towards Cpg dinucleotide sequences. Human 5-MCDG can be used
XX CC in a claimed method for labeling Cpg sequences corresponding to
XX CC methylated Cpg sequences in an isolated DNA sample. The method comprises:
XX CC digesting the genomic DNA with a restriction endonuclease to produce
XX CC genomic DNA fragments; treating the genomic DNA fragments with 5-MCDG
XX CC such that one or more 5-methylcytosine bases are removed to produce
XX CC abasic genomic DNA fragments; and treating these abasic genomic DNA
XX CC fragments with base excision/repair enzymes in the presence of labelled
XX CC dCTP such that 5-methylcytosine removed from the genomic DNA fragments by
XX CC 5-MCDG is replaced by labelled cytosine in the one or more corresponding
XX CC positions of the abasic genomic DNA fragments to produce labelled genomic
XX CC DNA fragments, so that specific labelling of Cpg sequences corresponding
XX CC to methylated Cpg sequences is achieved. The 5-MCDG is also used in a
XX CC claimed method for comparing Cpg methylation status, extent or pattern
XX CC between or among reference and test genomic DNA samples, and in a claimed
XX CC method for labelling potentially-methylatable Cpg sequences in Cpg-
XX CC containing genomic DNA fragments. The methods are used to identify
XX CC methylated and/or potentially methylatable Cpg dinucleotides in genomic
XX CC DNA, including comparison of methylation pattern between healthy and
XX CC diseased samples, for diagnosis.
XX
XX Sequence 307 AA;
SQ

Query Match 53.6%; Score 1639; DB 7; Length 307;
Best Local Similarity 100.0%; Pred. No. 8, 4e-150;
Matches 307; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 274 ESEPVAKSGOLDRTVCISDAGAGCETLSVTSENSLVKKKERLSGSGSNFCSEOKTSGII 333
|||||

DB 1 ESEPVAKSGOLDRTVCISDAGAGCETLSVTSENSLVKKKERLSGSGSNFCSEOKTSGII 60
QY 334 NKFCSAKDSSEHNKEKEDTFLESEBEGTGVYVERKEHATHTIILKRGSEMDNNGSPTRKDF 393
DB 61 NKFCSAKDSSEHNKEKEDTFLESEBEGTGVYVERKEHATHTIILKRGSEMDNNGSPTRKDF 120
QY 394 TGEKTFQEDDTLPRQIERRKTSLYFSSKYNEALSPPRKKAFFKMTTPRSPFNLYQETLFF 453
DB 121 TGEKTFQEDDTLPRQIERRKTSLYFSSKYNEALSPPRKKAFFKMTTPRSPFNLYQETLFF 180
QY 454 HDPWLLIATITFLNRTSGKMAIPVLWKFLEKYPSEAVARJADWRDVSLLKPLGLYDURA 513
DB 161 HDPWLLIATITFLNRTSGKMAIPVLWKFLEKYPSEAVARJADWRDVSLLKPLGLYDURA 240
QY 514 KTIKFSDEYLTQWKYPLELHGIGKYGNDSYRIFCVNEMQVHEDHKLNKYHDLWLEN 573
DB 241 KTIKFSDEYLTQWKYPLELHGIGKYGNDSYRIFCVNEMQVHEDHKLNKYHDLWLEN 300
QY 574 HEKLSLS 580
DB 301 HEKLSLS 307

RESULT 7
AAW88701
ID AAW88701 standard; protein; 257 AA.
XX
XX AAW88701;
AC
XX
XX 01-MAR-1999 (first entry)
XX
XX
XX Secreted protein encoded by gene 168 clone HCFNFI1.
XX
XX Human; secreted protein; fusion protein; gene therapy; protein therapy;
XX KW diagnosis; cissue; cancer; tumour; neurodegenerative disorder; leukaemia;
XX KW developmental abnormality; foetal deficiency; blood; allergy; renal;
XX KW immune system; aetna; lymphocytic disease; brain; hepatic; lymphoma;
XX KW inflammation; ischaemic shock; Alzheimer's disease; restenosis; AIDS;
XX KW cognitive disorder; schizophrenia; prostate; obesity; osteoclast; thymus;
XX KW osteoporosis; arthritis; testis; lung; thyroiditis; thyroid; digestion;
XX KW endocrine; metabolism; regulation; malabsorption; gastritis; neoplasm.
XX
XX Homo sapiens.
XX
XX WO9854963-A2.
XX
XX 10-DEC-1998.
XX
XX PD
XX
XX 04-JUN-1998; 98WO-US011422.
XX
XX 06-JUN-1997; 97US-0048875P.
XX PR 06-JUN-1997; 97US-0048876P.
XX PR 06-JUN-1997; 97US-0048877P.
XX PR 06-JUN-1997; 97US-0048878P.
XX PR 06-JUN-1997; 97US-0048880P.
XX PR 06-JUN-1997; 97US-0048881P.
XX PR 06-JUN-1997; 97US-0048882P.
XX PR 06-JUN-1997; 97US-0048883P.
XX PR 06-JUN-1997; 97US-0048884P.
XX PR 06-JUN-1997; 97US-0048885P.
XX PR 06-JUN-1997; 97US-0048892P.
XX PR 06-JUN-1997; 97US-0048893P.
XX PR 06-JUN-1997; 97US-0048894P.
XX PR 06-JUN-1997; 97US-0048895P.
XX PR 06-JUN-1997; 97US-0048896P.
XX PR 06-JUN-1997; 97US-0048897P.
XX PR 06-JUN-1997; 97US-0048898P.
XX PR 06-JUN-1997; 97US-0048899P.
XX PR 06-JUN-1997; 97US-0048900P.
XX PR 06-JUN-1997; 97US-0048901P.
XX PR 06-JUN-1997; 97US-0048915P.
XX PR 06-JUN-1997; 97US-0048916P.
XX PR 06-JUN-1997; 97US-0048917P.

PR 06-JUN-1997; 97US-0048949P.
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 PR 06-JUN-1997; 97US-0048970P.
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 PR 06-JUN-1997; 97US-0048974P.
 PR 06-JUN-1997; 97US-0049019P.
 PR 06-JUN-1997; 97US-0049020P.
 PR 06-JUN-1997; 97US-0049373P.
 PR 06-JUN-1997; 97US-0049374P.
 PR 06-JUN-1997; 97US-0049375P.
 PR 05-SEP-1997; 97US-0051584P.
 PR 05-SEP-1997; 97US-0051627P.
 PR 05-SEP-1997; 97US-0051628P.
 PR 05-SEP-1997; 97US-0051629P.
 PR 05-SEP-1997; 97US-0051634P.
 PR 05-SEP-1997; 97US-0051635P.
 PR 05-SEP-1997; 97US-0051642P.
 PR 05-SEP-1997; 97US-0051643P.
 PR 05-SEP-1997; 97US-0051644P.
 PR 05-SEP-1997; 97US-0051645P.
 PR 05-SEP-1997; 97US-0051646P.
 PR 05-SEP-1997; 97US-0051647P.
 PR 05-SEP-1997; 97US-0051648P.
 PR 05-SEP-1997; 97US-0051649P.
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 PR 05-SEP-1997; 97US-0051651P.
 PR 05-SEP-1997; 97US-0051654P.
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 PR 05-SEP-1997; 97US-0051667P.
 PR 05-SEP-1997; 97US-0051668P.
 PR 05-SEP-1997; 97US-0051760P.
 PR 05-SEP-1997; 97US-0051761P.
 PR 05-SEP-1997; 97US-0051762P.
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 PR 05-SEP-1997; 97US-0051764P.
 PR 05-SEP-1997; 97US-0051765P.
 PR 05-SEP-1997; 97US-0051769P.
 PR 05-SEP-1997; 97US-0051770P.
 PR 05-SEP-1997; 97US-0051771P.
 PR 05-SEP-1997; 97US-0051774P.
 PR 05-SEP-1997; 97US-0051775P.
 PR 05-SEP-1997; 97US-0051776P.
 PR 05-SEP-1997; 97US-0051777P.
 PR 05-SEP-1997; 97US-0051778P.
 PR 18-DEC-1997; 97US-0070923P.
 XX
 PA (HUMA-) HUMAN GENOME SCT INC.
 XX
 PI Young P, Greene JM, Ferrie AM, Ruben SM, Rosen CA, Hu J;
 PI Olsen HS, Ebner R, Brewer LA, Moore PA, Shi Y, Florence C;
 PI Florence K, Lafleur DM, Ni J, Fan P, Fischer CL, Soppet DR;
 PI Li Y, Zeng Z, Kyaw H, Yu G, Feng P, Dillon PJ, Endress GA,
 PI Carter KC;
 XX
 XX WPI, 1999-059865/05.
 DR N-PSDB; AAV84578.
 XX
 PT New isolated human gene and the secreted polypeptides they encode -
 PT useful for diagnosis and treatment of e.g. cancer, neurological
 PT disorders, immune diseases, inflammation or blood disorders.
 XX
 XX Claim 11; Page 564-565; 772pp; English.
 CC
 CC The invention relates to nucleic acid sequences (AAV84411 to AAV84633)
 CC encoding human secreted proteins (AAV88534 to AAV88756). The secreted
 CC protein gene sequences are deposited with the ATCC under deposit numbers
 CC ATCC 97979, 97974, 97975, 97976, 97977, 209007, 209008, 209009, 209010,
 CC 209011, 209080, 209081, 209082, 209083, 209084, 209085, 209511. Host

CC cells comprising recombinant vectors containing the nucleic acid
 CC sequences are used for the recombinant production of the secreted
 CC proteins. The polynucleotide and amino acid sequences are useful for are
 CC useful for preventing, treating or ameliorating medical conditions e.g.
 CC by protein or gene therapy. Pathological conditions can be also diagnosed
 CC by determining the amount of the new polypeptides in a sample or by
 CC determining the presence of mutations in the new polynucleotides.
 CC Specific uses are described for each of the polynucleotides, based on
 CC which tissues they are most highly expressed in, and include developing
 CC products for the diagnosis or treatment of cancer, neurodegenerative
 CC disorders, developmental abnormalities and foetal deficiencies, blood
 CC disorders, tumours, leukemias, diseases of the immune system, autoimmune
 CC diseases, hepatic and renal disease, lymphomas, inflammation, allergies,
 CC ischemic shock, Alzheimer's and cognitive disorders, schizophrenia,
 CC osteoporosis, prostate diseases, obesity, disorders involving osteoclasts
 CC such as osteoporosis, arthritis or malignancies, diseases of testes, lung
 CC or thyroid, digestive/endocrine disorders, infections and AIDS. The
 CC polypeptides are also useful for identifying their binding partners. The
 CC present sequence represents human secreted protein (see descriptor line
 CC for gene number and clone identification)
 XX
 SQ Sequence 257 AA;
 Query Match 40.3%; Score 1230; DB 2; Length 257;
 Best local Similarity 93.1%; Pred. No. 3.3e-110;
 Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;
 QY 160 MAALSHLQNSNNNNLNRCKCKKDVMPSSSELSQNSGLNFTSTHLLKDEG 219
 Db 1 MAALSHLQNSNNNNLNRCKCKKDVMPSSSELSQNSGLNFTSTHLLKDEG 60
 QY 220 VDDVNFPRKVRKPKGVITLKGIPKTKKCKRSCSGFVQSDSKRESVCKADAESEPPVA 279
 Db 61 VDDVNFPRKVRKPKGVITLKGIPKTKKCKRSCSGFVQSDSKRESVCKADAESEPPVA 120
 QY 280 QKSQIDRTVCISDAGAGCEFTLSVTSENSLVKKKERSLSSGNSPFCSQTKTSGIINKCSA 339
 Db 121 QKSQIDRTVCISDAGAGCEFTLSVTSENSLVKKKERSLSSGNSPFCSQTKTSGIINKCSA 180
 QY 340 KDSHNKKEVDTPFESEIEGTQVVERKHEHLHTDILKRESDNNNSPFRKPTGEKIF 399
 Db 181 KDSHNKKEVDTPFESEIEGTQVVERKHEHLHTDILKRESDNNNSPFRKPTGEKIF 235
 QY 400 QEDTTPRTQIERKRTSLYF 418
 Db 236 -EDTTPRTQIERKRTSLYF 253
 RESULT 8
 ABB50468
 ID ABB50468 standard; protein; 257 AA.
 XX
 AC ABB50468;
 XX
 DT 07-FEB-2002 (first entry)
 XX
 DE Human secreted protein encoded by gene 168 SEQ ID NO:416.
 XX
 KW Human; secreted protein; immunomodulatory; antisclerotic; anti-HIV;
 KW dermatological; immunosuppressive; anti-inflammatory; immunostimulant;
 KW cytoskeletal; cardiac; vascular; anti-angiogenic; ophthalmological;
 KW neuroprotective; nootropic; anticonvulsant; antialzheimers; vulnerary;
 KW antiparkinsonian; antimicrobial; gene therapy; vaccine; immune disorder;
 KW multiple sclerosis; systemic lupus erythematosus; HIV infection; cancer;
 KW human immunodeficiency virus; hyperproliferative disorder; wound healing;
 KW Gaucher's disease; cardiovascular disease; Schmidt's syndrome; chemotaxis;
 KW Chagas' cardiomyopathy; coronary arteriosclerosis; angiotensin disorder;
 KW corneal graft neovascularization; diabetic retinopathy; regeneration;
 KW neurological disorder; Huntington's chorea; Alzheimer's disease;
 KW Parkinson's disease; infectious disease; chromosome 3.
 KW
 KW Homo sapiens.
 XX
 OS

PN WO200162891-A2.
XX
XX 30-AUG-2001.
XX
XX 21-FEB-2001; 2001WO-US005614.
XX
XX 24-FEB-2000; 2000US-0184836P.
XX 29-MAR-2000; 2000US-0193170P.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Ni J, Ebner R, Lafleur DM, Moore PA, Olsen HS, Rosen CA;
PI Ruben SM, Soppet DR, Young PE, Shi Y, Florence KA, Mei Y;
PI Florence C, Hu J, Li Y, Kyaw H, Fischer CL, Ferris AM, Fan P;
PI Peng P, Andrews GA, Dillon PJ, Carter KC, Brewer LA, Yu G, Zeng Z,
PI Greene JM;
XX
XX WPI; 2001-625724/72.
DR N-PSDB; ABA83361.
XX
XX Nucleic acids encoding 207 human secreted polypeptides, useful for
PI preventing, diagnosing and/or treating, e.g. cancers, Parkinson's disease
PI and diabetic retinopathy.
XX
XX Claim 11, Page 1171-1172; 1533pp; English.
XX
XX ABB50301 to ABB51287 and ABA83194 to ABA83441 represent human secreted
CC proteins (I) and polynucleotide (II) sequences. (I) and (II) have various
CC activities based on the tissues and cells the genes are expressed in.
CC Example of these activities include: immunomodulatory; antisclerotic;
CC dermatological; immunosuppressive; anti-inflammatory; immunostimulant;
CC anti-HIV; cytostatic; cardiant; anti-angiogenic; ophthalmological;
CC neuroprotective; nootropic; anticonvulsant; antialzheimer's; vascular;
CC antiparkinsonian; antimicrobial; and vulnerary. (I) and (II) can be used
CC in gene therapy and vaccine production. (I) and (II) can be used in the
CC prevention, diagnosis and treatment of immune disorders (e.g. multiple
CC sclerosis, systemic lupus erythematosus and human immunodeficiency virus
CC (HIV) infections), hyperproliferative disorders (e.g. cancers and
CC Gaucher's disease), cardiovascular diseases (e.g. Schmitz syndrome,
CC Chaga's cardiomyopathy and coronary arteriosclerosis), angiogenic
CC disorders (e.g. corneal graft neovascularisation and diabetic
CC retinopathy), neurological disorders (e.g. Huntington's chorea,
CC Alzheimer's disease and Parkinson's disease), infectious diseases and/or
CC for promoting wound healing, regeneration and/or chemotaxis. ABA83185 to
CC ABA83193 and ABB50300 represent sequences used in the exemplification of
CC the present invention
XX
XX Sequence 257 AA;
SQ
Query Match 40.3%; Score 1230; DB 4; Length 257;
Best Local Similarity 93.1%; Pred. No. 3.3e-110;
Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;
QY 160 MAALSHLONQNSNMNRKTSKCKKQVFMPPSSSELOEGRGINSFTSHLLKEDG 219
DB 1 MAALSHLONQNSNMNRKTSKCKKQVFMPPSSSELOEGRGINSFTSHLLKEDG 60
QY 220 VDVNFRKRRKPKGKVTILKGIPIKTKKGRKSCGFPQSDSKESVCKKADASEPVA 279
DB 61 VDVNFRKRRKPKGKVTILKGIPIKTKKGRKSCGFPQSDSKESVCKKADASEPVA 120
QY 280 QKSOLDRTVCISDAGAGCETTLVTSSENSLVKKKERSLSSGSNFCSEQKTSGLINKFCGA 339
DB 121 QKSOLDRTVCISDAGAGCETTLVTSSENSLVKKKERSLSSGSNFCSEQKTSGLINKFCGA 180
QY 340 KQSEHNKEKEDPTLSESEIGTKVVERKEHLATDILKXGSEMDNCSPTKDFGEKTF 399
DB 181 KQSEHNKEKEDPTLSESEIGTKVVERKEHLATDILKXGSEMDNCSPTKDFGEKTF 235
QY 400 QEDTTPRTQIERRKTSLYF 418
DB 236 -EDTTPRNTDRKKENKPVF 253

RESULT 9
ABO44725
ID ABO44725 standard; protein: 257 AA.
XX
XX ABO44725;
XX
XX 02-OCT-2003 (first entry)
XX
XX Novel human secreted protein #168.
XX
XX Human; gene therapy; autoimmune disorder; multiple sclerosis; cancer;
XX systemic lupus erythematosus; haematopoietic cell disorder; allergy;
XX agammaglobulinaemia; ataxia telangiectasia; blood coagulation disorder;
XX athypogonadism; thrombocytopenia; graft-versus-host disease; arthritis;
XX inflammatory condition; ischaemia-reperfusion injury; infectious disease;
XX hyperproliferative disorder; purpura; viral infection; regeneration;
XX bacterial infection; ulcer; Alzheimer's disease.
XX
XX Homo sapiens.
XX
XX US2003065160-A1.
XX
XX 03-APR-2003.
XX
XX 07-DEC-2001; 2001US-00004860.
XX
XX 06-JUN-1997; 97US-0048875P.
XX 06-JUN-1997; 97US-0048876P.
XX 06-JUN-1997; 97US-0048877P.
XX 06-JUN-1997; 97US-0048878P.
XX 06-JUN-1997; 97US-0048880P.
XX 06-JUN-1997; 97US-0048881P.
XX 06-JUN-1997; 97US-0048882P.
XX 06-JUN-1997; 97US-0048883P.
XX 06-JUN-1997; 97US-0048884P.
XX 06-JUN-1997; 97US-0048892P.
XX 06-JUN-1997; 97US-0048893P.
XX 06-JUN-1997; 97US-0048894P.
XX 06-JUN-1997; 97US-0048895P.
XX 06-JUN-1997; 97US-0048896P.
XX 06-JUN-1997; 97US-0048897P.
XX 06-JUN-1997; 97US-0048898P.
XX 06-JUN-1997; 97US-0048899P.
XX 06-JUN-1997; 97US-0048900P.
XX 06-JUN-1997; 97US-0048901P.
XX 06-JUN-1997; 97US-0048915P.
XX 06-JUN-1997; 97US-0048916P.
XX 06-JUN-1997; 97US-0048917P.
XX 06-JUN-1997; 97US-0048949P.
XX 06-JUN-1997; 97US-0048962P.
XX 06-JUN-1997; 97US-0048963P.
XX 06-JUN-1997; 97US-0048964P.
XX 06-JUN-1997; 97US-0048970P.
XX 06-JUN-1997; 97US-0048971P.
XX 06-JUN-1997; 97US-0048972P.
XX 06-JUN-1997; 97US-0048974P.
XX 06-JUN-1997; 97US-0049019P.
XX 06-JUN-1997; 97US-0049020P.
XX 06-JUN-1997; 97US-0049373P.
XX 06-JUN-1997; 97US-0049374P.
XX 06-JUN-1997; 97US-0049375P.
XX 05-SEP-1997; 97US-0051584P.
XX 05-SEP-1997; 97US-0051627P.
XX 05-SEP-1997; 97US-0051628P.
XX 05-SEP-1997; 97US-0051634P.
XX 05-SEP-1997; 97US-0051635P.
XX 05-SEP-1997; 97US-0051642P.
XX 05-SEP-1997; 97US-0051643P.
XX 05-SEP-1997; 97US-0051644P.
XX 05-SEP-1997; 97US-0051645P.

Accession	Protein Name	Length	Score	E-value	Identity	Positives	Gaps	Conserved Domains
QY	220 VDVVNRKRRKRGKQKTTILKGIPIKTKKKGCKSCSGVQSDSKRESEVCKKADAESPVA	279						
Db	61 VDVVNRKRRKRGKQKTTILKGIPIKTKKKGCKSCSGVQSDSKRESEVCKKADAESPVA	120						
QY	280 QKSQLRATVCISDAGACGETTSLVSTSEENSVLVKKKERSLSSGSNFCSEQKTSGLINKFCSA	339						
Db	121 QKSQLRATVCISDAGACGETTSLVSTSEENSVLVKKKERSLSSGSNFCSEQKTSGLINKFCSA	180						
QY	340 KQSEHNKYEADTFLESEBIGTVKEVEVERKEHLATDILKRGSEMDNNCSPIRKDPTGEEKIF	399						
Db	181 KQSEHNKYEADTFLESEBIGTVKEVEVERKEHLATDILKRGSEMDNNCSPIRKDPTGEEKIF	235						
QY	400 QEDTIPRTQIERKTSLYF 418							
Db	236 -EDTIPRNTDRKKENKPV 253							
RESULT 10								
AB026205	AB026205 standard; protein; 257 AA.							
XX	AB026205;							
XX	10-SEP-2003 (first entry)							
XX	Human protein from novel secreted protein gene 160.							
XX	Human; secreted protein; precerebellin-like protein; neurodegenerative disorder; behavioural disorder; Alzheimer's disease; Parkinson's disease; Huntington's disease; schizophrenia; mania; dementia; paranoia; psychosis; autism; immune disorder; infection; inflammation; allergy; liver disorder; hepatoblastoma; jaundice; hepatitis; immunological disorder; AIDS; leukaemia; rheumatoid arthritis; sepsis; acne; psoriasis; cancer.							
XX	Homo sapiens.							
XX	US6525174-B1.							
XX	25-FEB-2003.							
PD	04-DEC-1998; 98US-00205258.							
XX	06-JUN-1997; 97US-0048875P.							
PR	06-JUN-1997; 97US-0048876P.							
PR	06-JUN-1997; 97US-0048877P.							
PR	06-JUN-1997; 97US-0048878P.							
PR	06-JUN-1997; 97US-0048880P.							
PR	06-JUN-1997; 97US-0048881P.							
PR	06-JUN-1997; 97US-0048882P.							
PR	06-JUN-1997; 97US-0048883P.							
PR	06-JUN-1997; 97US-0048884P.							
PR	06-JUN-1997; 97US-0048885P.							
PR	06-JUN-1997; 97US-0048892P.							
PR	06-JUN-1997; 97US-0048893P.							
PR	06-JUN-1997; 97US-0048894P.							
PR	06-JUN-1997; 97US-0048895P.							
PR	06-JUN-1997; 97US-0048900P.							
PR	06-JUN-1997; 97US-0048901P.							
PR	06-JUN-1997; 97US-0048915P.							
PR	06-JUN-1997; 97US-0048916P.							
PR	06-JUN-1997; 97US-0048917P.							
PR	06-JUN-1997; 97US-0048949P.							
PR	06-JUN-1997; 97US-0048962P.							
PR	06-JUN-1997; 97US-0048963P.							
PR	06-JUN-1997; 97US-0048964P.							
PR	06-JUN-1997; 97US-0048970P.							
PR	06-JUN-1997; 97US-0048971P.							

PR 06-JUN-1997; 97US-0048972P.
 PR 06-JUN-1997; 97US-0048974P.
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 PR 06-JUN-1997; 97US-0049020P.
 PR 06-JUN-1997; 97US-0049373P.
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 PR 05-SEP-1997; 97US-0057666P.
 PR 05-SEP-1997; 97US-0057667P.
 PR 05-SEP-1997; 97US-0057668P.
 PR 05-SEP-1997; 97US-0057760P.
 PR 05-SEP-1997; 97US-0057761P.
 PR 05-SEP-1997; 97US-0057762P.
 PR 05-SEP-1997; 97US-0057763P.
 PR 05-SEP-1997; 97US-0057765P.
 PR 05-SEP-1997; 97US-0057766P.
 PR 05-SEP-1997; 97US-0057769P.
 PR 05-SEP-1997; 97US-0057770P.
 PR 05-SEP-1997; 97US-0057771P.
 PR 05-SEP-1997; 97US-0057774P.
 PR 05-SEP-1997; 97US-0057775P.
 PR 05-SEP-1997; 97US-0057776P.
 PR 05-SEP-1997; 97US-0057777P.
 PR 05-SEP-1997; 97US-0057778P.
 PR 18-DEC-1997; 97US-0070923P.
 PR 04-JUN-1998; 98WO-US011422.
 PR 15-JUL-1998; 98US-0092921P.
 PR 30-JUL-1998; 98US-0094657P.
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PI Young P, Greene JM, Ferrie AM, Ruben SM, Rosen CA, Hu J,
 PI Olsen HS, Ehner R, Brewer LA, Moore PA, Shi Y, Florence C;
 PI Florence K, Lafleur DW, Ni J, Fan P, Wei Y, Fischer CL, Soppet DR,
 PI Li Y, Zeng Z, Kyaw H, Yu G, Feng P, Dillon PJ, Endress GA,
 PI Carter KC;
 XX
 DR WPI: 2003-511926/48.
 N-PSDB; ACD44672.
 XX
 PT New precerebellin-like protein, useful for diagnosing or treating
 PT neurodegenerative and behavioral disorders, immune disorders, liver
 PT disorders, and cancer.
 XX
 PS Disclosure; SEQ ID NO 416; 156bp; English.
 XX
 CC The invention relates to an isolated protein comprising amino acid
 CC residues 33-205 or 1-205 of a novel human secreted protein appearing as
 CC ABO26252. The protein is encoded by one of 238 disclosed cDNA sequences
 CC encoding 238 secreted proteins. ABO26252 is a precerebellin-like protein.
 CC Also included are a composition comprising the protein and a carrier and
 CC an isolated protein produced by expressing the protein cited above by a
 CC cell, and recovering the protein. The proteins are useful for diagnosing
 CC or treating neurodegenerative and behavioral disorders (e.g. Alzheimer's

CC disease, Parkinson's disease, Huntington's disease, schizophrenia, mania,
 CC dementia, paranoia, psychoses or autism), immune disorders (e.g.
 CC infection, inflammation, allergy), liver disorders (e.g. hepatoblastoma,
 CC jaundice, hepatitis), immunological disorders (e.g. AIDS, leukemia,
 CC rheumatoid arthritis, sepsis, acne, psoriasis) and cancer. The present
 CC sequence is one of the 238 disclosed novel secreted proteins. Note: The
 CC sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from USPTO
 CC at: seqdata.uspto.gov/sequence.html?docid=6525174B1
 XX
 SQ Sequence 257 AA;
 Query Match 40.3%; Score 1230; DB 7; Length 257;
 Best Local Similarity 93.1%; Pred. No. 3,3e-110;
 Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;
 Oy 160 MAALTSHLONSNNSNNLRTSRKCKDVFMPSSSSSLQSRGLSNTSTHLIKEDBG 219
 Db 1 MAALTSHLQNSNNNSNNLRTSRKCKDVFMPSSSSSLQSRGLSNTSTHLIKEDBG 60
 Oy 220 VDDVNFRRKRPKGVTLIKGPIPIKTKGCRKSCSGFVQSDSKRESCNADAESEPA 279
 Db 61 VDDVNFRRKRPKGVTLIKGPIPIKTKGCRKSCSGFVQSDSKRESCNADAESEPA 120
 Oy 280 QKSQLDRTVCISDAGAGETLSVTSSENSLVKKERSLSSGSNFCSEQKTSGLINKFCSA 339
 Db 121 QKSQLDRTVCISDAGAGETLSVTSSENSLVKKERSLSSGSNFCSEQKTSGLINKFCSA 180
 Oy 340 KDSHNEKYEEDTFLSESEIGTKVEVERKEHLHTDILKRGSEMDNCSPTKDTGKEIF 399
 Db 181 KDSHNEKYEEDTFLSESEIGTKVEVERKEHLHTDILKRGSEMDNCSPTKDTGKEIF 235
 Oy 400 QEDTIPRQIERKRTSLYF 418
 Db 236 -EDTIPRNTDRKKNKPVF 253
 RESULT 11
 ADD89916 standard; protein; 202 AA.
 ID ADD89916
 XX
 AC ADD89916;
 XX
 DT 29-JAN-2004 (first entry)
 XX
 DE Human 5-methylcytosine DNA glycosylase N-terminal deletion mutant.
 XX
 KW Human; 5-methylcytosine DNA glycosylase; enzyme; Cpg; mutant; mutcin.
 XX
 OS Homo sapiens.
 XX
 PN WO2003078593-A2.
 XX
 PD 25-SEP-2003.
 XX
 PF 14-MAR-2003; 2003WO-US007933.
 XX
 PR 15-MAR-2002; 2002US-0364689P.
 XX
 PA (EPIC-) EPIGENOMICS AG.
 XX
 PT Lofton-Day CE, Day JK;
 XX
 DR WPI: 2003-779127/73.
 XX
 PT Labeling methylated or methylatable Cpg sequences, useful e.g. for
 PT diagnostic detection of altered methylation, comprises replacing
 PT methylated cytosine by labeled cytosine.
 XX
 PS Claim 11; Page 71-72; 73pp; English.
 XX
 CC The present sequence is the protein sequence of an N-terminal deletion
 CC mutant of human 5-methylcytosine DNA glycosylase (5-MCG), in which amino

Query Match 11.7%; Score 357; DB 5; Length 68;
Best Local Similarity 98.5%; Pred. No. 2.1e-26;
Matches 67; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 81 ECRSVPCGWRVYKORLFGTAGRFDYFTSPQGLKFRSSSLANYLHKNGETSLKPED 140
DB 1 ECRSVPCGWRVYKORLFGTAGRFDYFTSPQGLKFRSSSLANYLHKNGETSLKPED 60

QY 141 FDFTVLSK 148
DB 61 FDFTVLSK 68

RESULT 15
ABB51147
ID ABB51147 standard; protein; 50 AA.
XX ABB51147;
AC 07-FEB-2002 (first entry)
DT 07-FEB-2002 (first entry)
XX Human secreted protein encoded by gene 168 SEQ ID NO:1100.
DE Human, secreted protein; immunomodulatory; antisclerotic; anti-HIV;
XX dermatological; immunosuppressive; antiinflammatory; immunostimulant;
KW cyostatic; cardiant; vascular; anti-angiogenic; ophthalmological;
KW neuroprotective; noctropic; anticonvulsant; antialzheimers; vulnerary;
KW antiparkinsonian; antitremoral; gene therapy; vaccine; immune disorder;
KW multiple sclerosis; systemic lupus erythematosus; HIV infection; cancer;
KW human immunodeficiency virus; hyperproliferative disorder; wound healing;
KW Gaucher's disease; cardiovascular disease; Schmitz syndrome; chemotaxis;
KW Chaga's cardiomyopathy; coronary arteriosclerosis; angio-genic disorder;
KW corneal graft neovascularization; diabetic retinopathy; regeneration;
KW neurological disorder; Huntington's chorea; Alzheimer's disease;
KW Parkinson's disease; infectious disease; chromosome 3.
XX Homo sapiens.
OS WO200162891-A2.
XX 30-AUG-2001.
PD 21-FEB-2001; 2001WO-US005614.
XX 24-FEB-2000; 2000US-0184836P.
PR 29-MAR-2000; 2000US-0193170P.
XX (HUMA-) HUMAN GENOME SCI INC.
PA Ni J, Ebner R, Lafleur DW, Moore PA, Olsen HS, Rosen CA;
XX Ruben SM, Soppet DR, Young PE, Shi Y, Florence KA, Wei Y;
PI Florence C, Hu J, Li Y, Kyaw H, Fischer CJ, Ferrie AM, Fan P;
PI Feng P, Endress GA, Dillon PJ, Carter KC, Brewer LA, Yu G, Zeng Z;
PI Greene JM;
XX WPI; 2001-625724/72.
DR Nucleic acids encoding 207 human secreted polypeptides, useful for
XX preventing, diagnosing and/or treating, e.g. cancers, Parkinson's disease
PT and diabetic retinopathy.
PT Disclosure; Page 366; 1533pp; English.
PS ABB50301 to ABB51287 and ABA83194 to ABA83441 represent human secreted
XX proteins (I) and polynucleotides (II) sequences. (I) and (II) have various
CC activities based on the tissues and cells the genes are expressed in.
CC Example of these activities include: immunomodulatory; antisclerotic;
CC dermatological; immunosuppressive; antiinflammatory; immunostimulant;
CC anti-HIV; cyostatic; cardiant; anti-angiogenic; ophthalmological;
CC neuroprotective; noctropic; anticonvulsant; antialzheimers; vascular;
CC antiparkinsonian; antitremoral; and vulnerary. (I) and (II) can be used
CC in gene therapy and vaccine production. (I) and (II) can be used in the
CC prevention, diagnosis and treatment of immune disorders (e.g. multiple

CC sclerosis, systemic lupus erythematosus and human immunodeficiency virus
CC (HIV) infections), hyperproliferative disorders (e.g. cancers and
CC Gaucher's disease), cardiovascular diseases (e.g. Schmitz syndrome,
CC Chaga's cardiomyopathy and coronary arteriosclerosis), angiogenic
CC disorders (e.g. corneal graft neovascularisation and diabetic
CC retinopathy), neurological disorders (e.g. Huntington's chorea,
CC Alzheimer's disease and Parkinson's disease), infectious diseases and/or
CC for promoting wound healing, regeneration and/or chemotaxis. ABA83185 to
CC ABA83193 and ABB50300 represent sequences used in the exemplification of
CC the present invention
XX Sequence 50 AA;
SQ

Query Match 8.3%; Score 254; DB 4; Length 50;
Best Local Similarity 98.0%; Pred. No. 1.3e-16;
Matches 49; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 110 FISPQGLKFRSSSLANYLHKNGETSLKPEDFDTVLSKRGISRYDCS 159
DB 1 FISPQGLKFRSSSLANYLHKNGETSLKPEDFDTVLSKRGISRYDCS 50

Search completed: August 22, 2005, 10:05:09
Job time : 172 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 22, 2005, 10:02:09 ; Search time 163 Seconds
(without alignments)
1393.374 Million cell updates/sec

Title: US-10-629-951-2

Perfect score: 3055
Sequence: 1 MGTGLSLSLDGGAAPTV.....HKLNKTHDMLWENHEKLSLS 580

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1759131 seqs, 391586102 residues

Total number of hits satisfying chosen parameters: 1759131

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

1: /cgn2_6/ptodata/1/pubppaa/US07_PUBCOMB.pep.*
2: /cgn2_6/ptodata/1/pubppaa/PCT_NEW_PUB.pep.*
3: /cgn2_6/ptodata/1/pubppaa/US06_NEW_PUB.pep.*
4: /cgn2_6/ptodata/1/pubppaa/US06_PUBCOMB.pep.*
5: /cgn2_6/ptodata/1/pubppaa/US07_NEW_PUB.pep.*
6: /cgn2_6/ptodata/1/pubppaa/PCTUS_PUBCOMB.pep.*
7: /cgn2_6/ptodata/1/pubppaa/US08_NEW_PUB.pep.*
8: /cgn2_6/ptodata/1/pubppaa/US08_PUBCOMB.pep.*
9: /cgn2_6/ptodata/1/pubppaa/US09_PUBCOMB.pep.*
10: /cgn2_6/ptodata/1/pubppaa/US09_PUBCOMB.pep.*
11: /cgn2_6/ptodata/1/pubppaa/US09_PUBCOMB.pep.*
12: /cgn2_6/ptodata/1/pubppaa/US10_NEW_PUB.pep.*
13: /cgn2_6/ptodata/1/pubppaa/US10_PUBCOMB.pep.*
14: /cgn2_6/ptodata/1/pubppaa/US10_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubppaa/US10_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubppaa/US10_PUBCOMB.pep.*
17: /cgn2_6/ptodata/1/pubppaa/US10_PUBCOMB.pep.*
18: /cgn2_6/ptodata/1/pubppaa/US10_NEW_PUB.pep.*
19: /cgn2_6/ptodata/1/pubppaa/US11_PUBCOMB.pep.*
20: /cgn2_6/ptodata/1/pubppaa/US11_NEW_PUB.pep.*
21: /cgn2_6/ptodata/1/pubppaa/US60_NEW_PUB.pep.*
22: /cgn2_6/ptodata/1/pubppaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	3055	100.0	580	US-10-389-853-2	Sequence 2, Appli
2	3055	100.0	580	US-10-629-951-2	Sequence 2, Appli
3	3011	98.6	574	US-10-629-951-24	Sequence 24, Appl
4	1639	53.6	307	US-10-389-853-11	Sequence 11, Appl
5	1326	43.4	384	US-10-629-951-29	Sequence 29, Appl
6	1230	40.3	257	US-09-933-767-416	Sequence 416, App
7	1230	40.3	257	US-10-004-860-416	Sequence 416, App
8	1230	40.3	257	US-10-023-282-416	Sequence 416, App
9	1106	36.2	202	US-10-389-853-12	Sequence 12, Appl
10	880.5	28.8	416	US-10-389-853-4	Sequence 4, Appli
11	816	26.7	147	US-10-389-853-13	Sequence 13, Appl

12	697	22.8	126	15	US-10-629-951-32	Sequence 32, Appl
13	616	20.2	119	15	US-10-629-951-30	Sequence 30, Appl
14	443	14.5	85	15	US-10-629-951-37	Sequence 37, Appl
15	357	11.7	68	10	US-09-967-869A-7	Sequence 7, Appli
16	357	11.7	68	20	US-11-045-828-7	Sequence 7, Appli
17	330	10.8	443	16	US-10-739-930-5943	Sequence 5943, Ap
18	300.5	9.8	382	16	US-10-425-115-299452	Sequence 299452,
19	281	9.2	185	16	US-10-767-701-33975	Sequence 33975, A
20	254	8.3	50	10	US-09-933-767-1100	Sequence 1100, Ap
21	254	8.3	50	14	US-10-004-860-1100	Sequence 1100, Ap
22	254	8.3	50	14	US-10-023-282-1100	Sequence 1100, Ap
23	209	6.8	476	16	US-10-408-765A-266	Sequence 266, Appl
24	208.5	6.8	560	16	US-10-475-681-7	Sequence 7, Appli
25	208.5	6.8	561	16	US-10-475-681-3	Sequence 31, Appli
26	192	6.1	132	15	US-10-629-951-31	Sequence 31, Appli
27	186	6.1	68	10	US-09-967-869A-5	Sequence 5, Appli
28	186	6.1	68	20	US-11-045-828-5	Sequence 5, Appli
29	152	5.0	105	16	US-10-437-963-148365	Sequence 148365,
30	145	4.7	1301	15	US-10-369-493-1644	Sequence 1644, Ap
31	145	4.7	3418	15	US-10-392-113-42	Sequence 42, Appli
32	145	4.7	3418	15	US-10-634-574-1	Sequence 1, Appli
33	145	4.7	3418	16	US-10-408-765A-178	Sequence 178, App
34	137	4.5	560	14	US-10-205-841-12	Sequence 12, Appl
35	137	4.5	1713	18	US-10-840-512-116	Sequence 116, App
36	136.5	4.5	409	16	US-10-425-115-250086	Sequence 250086,
37	136.5	4.5	1359	17	US-10-732-923-8708	Sequence 8708, Ap
38	135.5	4.4	282	10	US-09-967-869A-17	Sequence 17, Appl
39	135.5	4.4	282	20	US-11-045-828-17	Sequence 17, Appl
40	135.5	4.4	755	16	US-10-473-127-548	Sequence 548, App
41	135.5	4.4	755	16	US-10-473-127-550	Sequence 550, App
42	135.5	4.4	755	16	US-10-473-127-553	Sequence 553, App
43	133.5	4.4	1359	17	US-10-732-923-8707	Sequence 8707, Ap
44	132	4.3	578	9	US-09-821-835-2	Sequence 2, Appli
45	132	4.3	1819	15	US-10-335-977-7981	Sequence 7981, Ap

ALIGNMENTS

RESULT 1
US-10-389-853-2
1: Sequence 2, Application US//10389853
2: Publication No. US20030180779A1
GENERAL INFORMATION:
APPLICANT: Loftron-Day, Cathy E.
TITLE OF INVENTION: Diagnostic Methods Using 5-Methylcytosine DNA Glyco
FILE REFERENCE: 47675-36
CURRENT FILING DATE: 2003-03-14
CURRENT APPLICATION NUMBER: US/10/389, 853
PRIOR FILING DATE: 2002-03-14
PRIOR APPLICATION NUMBER: 60/364,689
NUMBER OF SEQ ID NOS: 13
SOFTWARE: PatentIn version 3.1
SEQ ID NO 2
LENGTH: 580
TYPE: PRT
ORGANISM: Homo sapiens
US-10-389-853-2

Query Match	100.0%	Score 3055	DB 14	Length 580
Best Local Similarity	100.0%	Pred. No. 8.2e-236		
Matches	580	Conservative	0	Mismatches 0; Indels 0; Gaps 0
QY	1	MGTGLSLSLDGGAAPTVSSERLVDPNDLRKEDVAMELERGEDEBQMIRKSSSE	60	
DB	1	MGTGLSLSLDGGAAPTVSSERLVDPNDLRKEDVAMELERGEDEBQMIRKSSSE	60	
QY	61	CNPLQEPYASQFGATAGTCRKSVCPCWERYVQRLFGKTRAGRPDYVFIPQGIKFS	120	
DB	61	CNPLQEPYASQFGATAGTCRKSVCPCWERYVQRLFGKTRAGRPDYVFIPQGIKFS	120	
QY	121	KSLIANYLHKNETSLKRPEDFTVLSKRGISRYDCSMALVTSHLQNSNNSMNLRT	180	

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Db 121 KSSLANTYHKNGETSLKPEDDFTVLSKRGIKSRKYDCSMAALTSHLQONSNNSNMNLT 180
Qy 181 RSKCKKQVFMPPSSSSSELOESRGLSNFTSTHLLKEDGVDDVNFRRKVRKPKGVTLTKG 240
Db 181 RSKCKKQVFMPPSSSSSELOESRGLSNFTSTHLLKEDGVDDVNFRRKVRKPKGVTLTKG 240
Qy 241 IPKTKKGGCRKSCSGFVQSDSKRESVCKNKADESEPAOKSOLDRTVCISDAGACGCTL 300
Db 241 IPKTKKGGCRKSCSGFVQSDSKRESVCKNKADESEPAOKSOLDRTVCISDAGACGCTL 300
Qy 301 SVTSEENSLVKKKERSLSSGSNFCSEQKTGIIINKFCSAKOSEHNEKXEDTFLSESEIGT 360
Db 301 SVTSEENSLVKKKERSLSSGSNFCSEQKTGIIINKFCSAKOSEHNEKXEDTFLSESEIGT 360
Qy 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFTEGKIPOEDTIPRQIERRTSLYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFTEGKIPOEDTIPRQIERRTSLYFSS 420
Qy 421 KYNKEALSPRRKAFKMTPPRSPFNLVQETLPHDPKLLIATIFLNRTSGKMAIPVLMK 480
Db 421 KYNKEALSPRRKAFKMTPPRSPFNLVQETLPHDPKLLIATIFLNRTSGKMAIPVLMK 480
Qy 481 FLEKYPBAEVAARTADWDVSELKPLGLYDLRAKTIYKFSDEYLTQWKYPILHGIKGY 540
Db 481 FLEKYPBAEVAARTADWDVSELKPLGLYDLRAKTIYKFSDEYLTQWKYPILHGIKGY 540
Qy 541 GNDSYRIFCVNEMKQVAPEDHKLNTKYHDMLENHEKSLS 580
Db 541 GNDSYRIFCVNEMKQVAPEDHKLNTKYHDMLENHEKSLS 580
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RESULT 2

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US-10-629-951-2
; Sequence 2, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629,951
; PRIOR FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2
; LENGTH: 580
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-629-951-2
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Query Match 100.0%; Score 3055; DB 15; Length 580;
Best Local Similarity 100.0%; Pred. No. 8,2e-236;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Qy 1 MGTTLGSLSLGDRGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEEQMMIKRSSE 60
Db 1 MGTTLGSLSLGDRGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEEQMMIKRSSE 60
Qy 61 CNPLQEPPIASAOAGTAGTECRKSVPCGWERVVQKQLFGKTAGRFDVYFISPOGLKFRS 120
Db 61 CNPLQEPPIASAOAGTAGTECRKSVPCGWERVVQKQLFGKTAGRFDVYFISPOGLKFRS 120
Qy 121 KSSLANTYHKNGETSLKPEDDFTVLSKRGIKSRKYDCSMAALTSHLQONSNNSNMNLT 180
Db 121 KSSLANTYHKNGETSLKPEDDFTVLSKRGIKSRKYDCSMAALTSHLQONSNNSNMNLT 180
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Db 121 KSSLANTYHKNGETSLKPEDDFTVLSKRGIKSRKYDCSMAALTSHLQONSNNSNMNLT 180
Qy 181 RSKCKKQVFMPPSSSSSELOESRGLSNFTSTHLLKEDGVDDVNFRRKVRKPKGVTLTKG 240
Db 181 RSKCKKQVFMPPSSSSSELOESRGLSNFTSTHLLKEDGVDDVNFRRKVRKPKGVTLTKG 240
Qy 241 IPKTKKGGCRKSCSGFVQSDSKRESVCKNKADESEPAOKSOLDRTVCISDAGACGCTL 300
Db 241 IPKTKKGGCRKSCSGFVQSDSKRESVCKNKADESEPAOKSOLDRTVCISDAGACGCTL 300
Qy 301 SVTSEENSLVKKKERSLSSGSNFCSEQKTGIIINKFCSAKOSEHNEKXEDTFLSESEIGT 360
Db 301 SVTSEENSLVKKKERSLSSGSNFCSEQKTGIIINKFCSAKOSEHNEKXEDTFLSESEIGT 360
Qy 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFTEGKIPOEDTIPRQIERRTSLYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFTEGKIPOEDTIPRQIERRTSLYFSS 420
Qy 421 KYNKEALSPRRKAFKMTPPRSPFNLVQETLPHDPKLLIATIFLNRTSGKMAIPVLMK 480
Db 421 KYNKEALSPRRKAFKMTPPRSPFNLVQETLPHDPKLLIATIFLNRTSGKMAIPVLMK 480
Qy 481 FLEKYPBAEVAARTADWDVSELKPLGLYDLRAKTIYKFSDEYLTQWKYPILHGIKGY 540
Db 481 FLEKYPBAEVAARTADWDVSELKPLGLYDLRAKTIYKFSDEYLTQWKYPILHGIKGY 540
Qy 541 GNDSYRIFCVNEMKQVAPEDHKLNTKYHDMLENHEKSLS 580
Db 541 GNDSYRIFCVNEMKQVAPEDHKLNTKYHDMLENHEKSLS 580
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RESULT 3

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US-10-629-951-24
; Sequence 24, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629,951
; PRIOR FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 24
; LENGTH: 574
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-629-951-24
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Query Match 98.6%; Score 3011; DB 15; Length 574;
Best Local Similarity 99.0%; Pred. No. 2.7e-232;
Matches 574; Conservative 0; Mismatches 0; Indels 6; Gaps 1;
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Qy 1 MGTTLGSLSLGDRGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEEQMMIKRSSE 60
Db 1 MGTTLGSLSLGDRGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEEQMMIKRSSE 60
Qy 61 CNPLQEPPIASAOAGTAGTECRKSVPCGWERVVQKQLFGKTAGRFDVYFISPOGLKFRS 120
Db 61 CNPLQEPPIASAOAGTAGTECRKSVPCGWERVVQKQLFGKTAGRFDVYFISPOGLKFRS 120
Qy 121 KSSLANTYHKNGETSLKPEDDFTVLSKRGIKSRKYDCSMAALTSHLQONSNNSNMNLT 180
Db 121 KSSLANTYHKNGETSLKPEDDFTVLSKRGIKSRKYDCSMAALTSHLQONSNNSNMNLT 180
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QY 181 RSKCKDVMPSPSSSEILOSERGLSNFTSTHLLKEDSGVDVDFNRKRYKPKGKTTLLKG 240
DB 181 RSKCKDVMPSPSSSEILOSERGLSNFTSTHLLKEDSGVDVDFNRKRYKPKGKTTLLKG 240
QY 241 IPIKTKKGCRCSCGFPVSDSKRESVCNKADAESEPVAKQSLDRTVCISDAGCGEYL 300
DB 241 IPIKTKKGCRCSCGFPVSDSKRESVCNKADAESEPVAKQSLDRTVCISDAGCGEYL 300
QY 301 SVTSEENSLVKKKERSLSGSGNFCSEOKTSGIINKFCSAKXSEHNEKXEDTFLSEBEIGT 360
DB 301 SVTSEENSLVKKKERSLSGSGNFCSEOKTSGIINKFCSAKXSEHNEKXEDTFLSEBEIGT 360
QY 361 KVEVERKEHHTDILKRGSEMDNNSPFRKOPFTGEKIFQEDTIPRTOIERKRTSLYFS 420
DB 361 KVEVERKEHHTDILKRGSEMDNNSPFRKOPFTGEKIFQEDTIPRTOIERKRTSLYFS 414
QY 421 KYNKALSPRRKAPFKWTPRSPFNLVOETLPHDPWKLLIATIFLNRSGMAIPVLMK 480
DB 421 KYNKALSPRRKAPFKWTPRSPFNLVOETLPHDPWKLLIATIFLNRSGMAIPVLMK 474
QY 445 FLEKTPSAEVARADMDVSEILKPLGLYDLRAKTIKFSDEYLTQWKYPIELHGIGKY 540
DB 445 FLEKTPSAEVARADMDVSEILKPLGLYDLRAKTIKFSDEYLTQWKYPIELHGIGKY 534
QY 541 GNDYRIFCVNEMKQVHPEDHKLKNTYHDLWLNHEKLSLS 580
DB 541 GNDYRIFCVNEMKQVHPEDHKLKNTYHDLWLNHEKLSLS 574

RESULT 4
US-10-389-853-11
/ Sequence 11, Application US/10389853
/ Publication No. US2003018079A1
/ GENERAL INFORMATION:
/ APPLICANT: Lofton-Day, Cathy E.
/ APPLICANT: Day, John K.
/ TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glyco
/ FILE REFERENCE: 47675-36
/ CURRENT APPLICATION NUMBER: US/10/389,853
/ CURRENT FILING DATE: 2003-03-14
/ PRIOR FILING DATE: 2002-03-15
/ PRIOR APPLICATION NUMBER: 60/364,689
/ NUMBER OF SEQ ID NOS: 13
/ SOFTWARE: PatentIn version 3.1
/ SEQ ID NO 11
/ LENGTH: 307
/ TYPE: PRT
/ ORGANISM: Homo sapiens
/ FEATURE:
/ NAME/KEY: NON TER
/ LOCATION: (1)-(11)
/ OTHER INFORMATION: functional N-terminal deletion mutant of hMBD4 (SEQ ID NO:2); aa
/ OTHER INFORMATION: no.1 corresponds to aa no.274 of hMBD4; mutant shows enhanced d
/ OTHER INFORMATION: eglycosylase specificity towards CpG dinucleotide sequences; see
/ OTHER INFORMATION: Zhu et al. Nuc. Acid Res. 28:4157-4165, 2000.
US-10-389-853-11

Query Match 53.6%; Score 1639; DB 14; Length 307;
Best Local Similarity 100.0%; Pred. No. 1e-122;
Matches 307; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 274 ESEPVAKQSLDRTVCISDAGCGEYLSVTSEENSLVKKKERSLSGSGNFCSEOKTSGII 333
DB 1 ESEPVAKQSLDRTVCISDAGCGEYLSVTSEENSLVKKKERSLSGSGNFCSEOKTSGII 60
QY 334 NKFCSAKXSEHNEKXEDTFLSEBEIGTKVEVERKEHHTDILKRGSEMDNNSPFRKOP 393
DB 61 NKFCSAKXSEHNEKXEDTFLSEBEIGTKVEVERKEHHTDILKRGSEMDNNSPFRKOP 120
QY 394 TGEKIFQEDTIPRTOIERKRTSLYFSKYNKALSPRRKAPFKWTPRSPFNLVOETL 453
DB 121 TGEKIFQEDTIPRTOIERKRTSLYFSKYNKALSPRRKAPFKWTPRSPFNLVOETL 180
```

```
QY 454 HDPWKLLIATIFLNRSGMAIPVLMKFLKYPASAEVARTADMDVSEILKPLGLYDLRA 513
DB 181 HDPWKLLIATIFLNRSGMAIPVLMKFLKYPASAEVARTADMDVSEILKPLGLYDLRA 240
QY 514 KTIKFSDEYLTQWKYPIELHGIGKYGNDYRIFCVNEMKQVHPEDHKLKNTYHDLWLN 573
DB 241 KTIKFSDEYLTQWKYPIELHGIGKYGNDYRIFCVNEMKQVHPEDHKLKNTYHDLWLN 300
QY 574 HEKLSLS 580
DB 301 HEKLSLS 307

RESULT 5
US-10-629-951-29
/ Sequence 29, Application US/10629951
/ Publication No. US20040018550A1
/ GENERAL INFORMATION:
/ APPLICANT: Bellacosa, Alfonso
/ TITLE OF INVENTION: Methods for Detection of Transition
/ FILE REFERENCE: PCCC 96-21
/ CURRENT APPLICATION NUMBER: US/10/629,951
/ CURRENT FILING DATE: 2003-07-29
/ PRIOR APPLICATION NUMBER: US/09/629,222A
/ PRIOR FILING DATE: 2000-07-31
/ PRIOR APPLICATION NUMBER: 09/463,891
/ PRIOR FILING DATE: 2000-01-28
/ PRIOR APPLICATION NUMBER: PCT/US98/15828
/ PRIOR FILING DATE: 1998-07-28
/ PRIOR APPLICATION NUMBER: 60/053,936
/ PRIOR FILING DATE: 1997-07-28
/ SOFTWARE: FaatSEO for Windows Version 3.0
/ SEQ ID NO 29
/ LENGTH: 384
/ TYPE: PRT
/ ORGANISM: Mus musculus
US-10-629-951-29

Query Match 43.4%; Score 1326; DB 15; Length 384;
Best Local Similarity 54.3%; Pred. No. 1.7e-97;
Matches 283; Conservative 29; Mismatches 71; Indels 138; Gaps 4;

QY 36 KEDVAMELERVGEDEEQMIRKSSSECNPLQEPISAPGATAGTECKRSVPQGWERYVK 95
DB 1 KEDVAVGLGVGEDGKDLVI--SSERSLLOEPTAST--LSSTATTEGHKVPVCGWERYVK 57
QY 96 QRLFKTKGRPDVYRTISFQGLKFRKSSLANYLHNKGETSLKPEDPFTVLSKRGISKRY 155
DB 58 QRLSKRTKGPVYRTISFQGLKFRKRSILANLNGEFLKPEDPFTVLPKSGSINPGY 117
QY 156 KDCSMAALTSHLQONSSNNMNLARTRSCKKQDVFMPPSSSELQESRGLSNFTSTHLLK 215
DB 118 KQSLAALSLLOPNETDVSKQNLKTRSKKTKDVLPDSTSSPSSSGISNNSGACILLR 177
QY 216 EDEGVDDVNFRRKRYKPKGKVTLLKGIPIKTKKGCRCSCGFPVSDSKRESVCNKADAES 275
DB 178 EHRDLDQDVDESKRRRSKRVTVLKGTASOKTKQCKRSKLESTQNRKRAAS----- 228
QY 276 EPVAKQSLDRTVCISDAGCGEYLSVTSEENSLVKKKERSLSGSGNFCSEOKTSGIINK 335
DB 229 ----- 228
QY 336 FCSAKXSEHNEKXEDTFLSEBEIGTKVEVERKEHHTDILKRGSEMDNNSPFRKOPFG 395
DB 229 ----- 228
QY 396 EKIQEDTIPRTOIERKRTSLYFSKYNKALSPRRKAPFKWTPRSPFNLVOETLPHD 455
DB 229 -----EDSIPRTOVERKRTSLYFSKYNKALSPRRKAPFKWTPRSPFNLVOETLPHD 283
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Db      1 MAALTSHLQNSNSWNLRTSKCKDVFMPPSSSSSELOESRGLSNFTSTHLLKEDG 60
QY      220 VDVNFRKVRKPKGKVTLLKGIPIKTKKCGCRKSCGFPVQSDSKESVCKNKADESEPA 279
Db      61 VDVNFRKVRKPKGKVTLLKGIPIKTKKCGCRKSCGFPVQSDSKESVCKNKADESEPA 120
QY      280 QKSQLDRTVCISDAGACGETLSVTSEENSLVKKKERSLSSGNSFCSEQKTSGLINKFCSA 339
Db      121 QKSQLDRTVCISDAGACGETLSVTSEENSLVKKKERSLSSGNSFCSEQKTSGLINKFCSA 180
QY      340 KQSENEKEEDFTLSEEEIGTVEVVERKEHLHTDILKRGSEMDNCSPTRKDFTGEKIF 399
Db      181 KQSENEKEEDFTLSEEEIGTVEVVERKEHLHTDILKRGSEMDNCSPTRKDFT----- 235
QY      400 QEDTIPRTQIERRKTSLYF 418
Db      236 -EDTIPRTQIERRKTSLYF 253

```

```

RESULT 7
US-10-004-860-416
; Sequence 416, Application US/10004860
; Publication No. US20030065160A1
; GENERAL INFORMATION:
; APPLICANT: Young et al.
; TITLE OF INVENTION: 207 Human Secreted Proteins
; FILE REFERENCE: P2007P1
; CURRENT APPLICATION NUMBER: US/10/004,860
; CURRENT FILING DATE: 2001-12-07
; Prior Application removed - See File Wrapper or Palm
; NUMBER OF SEQ ID NOS: 1227
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 416
; LENGTH: 257
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: SITE
; LOCATION: (100)
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
US-10-004-860-416

```

```

Query Match      40.3%; Score 1230; DB 14; Length 257;
Best Local Similarity 93.1%; Pred. No. 4.6e-90;
Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;

QY      160 MAALTSHLQNSNSWNLRTSKCKDVFMPPSSSSSELOESRGLSNFTSTHLLKEDG 219
Db      1 MAALTSHLQNSNSWNLRTSKCKDVFMPPSSSSSELOESRGLSNFTSTHLLKEDG 60
QY      220 VDVNFRKVRKPKGKVTLLKGIPIKTKKCGCRKSCGFPVQSDSKESVCKNKADESEPA 279
Db      61 VDVNFRKVRKPKGKVTLLKGIPIKTKKCGCRKSCGFPVQSDSKESVCKNKADESEPA 120
QY      280 QKSQLDRTVCISDAGACGETLSVTSEENSLVKKKERSLSSGNSFCSEQKTSGLINKFCSA 339
Db      121 QKSQLDRTVCISDAGACGETLSVTSEENSLVKKKERSLSSGNSFCSEQKTSGLINKFCSA 180
QY      340 KQSENEKEEDFTLSEEEIGTVEVVERKEHLHTDILKRGSEMDNCSPTRKDFTGEKIF 399
Db      181 KQSENEKEEDFTLSEEEIGTVEVVERKEHLHTDILKRGSEMDNCSPTRKDFT----- 235
QY      400 QEDTIPRTQIERRKTSLYF 418
Db      236 -EDTIPRTQIERRKTSLYF 253

```

```

RESULT 8
US-10-023-282-416
; Sequence 416, Application US/10023282
; Publication No. US20030092893A1
; GENERAL INFORMATION:

```

```

; APPLICANT: Young et al.
; TITLE OF INVENTION: 207 Human Secreted Proteins
; FILE REFERENCE: P2007P1
; CURRENT APPLICATION NUMBER: US/10/023,282
; CURRENT FILING DATE: 2001-12-20
; EARLIER APPLICATION NUMBER: 09/205,258
; EARLIER FILING DATE: 1998-12-04
; EARLIER APPLICATION NUMBER: PCT/US98/11422
; EARLIER FILING DATE: 1998-06-04
; EARLIER APPLICATION NUMBER: 60/048,885
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,375
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,881
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,880
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,896
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,020
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,876
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,895
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,884
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,894
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,971
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,964
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,882
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,899
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,893
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,900
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,901
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,892
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,915
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,019
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,970
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,972
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,916
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,373
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,875
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,374
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,917
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,949
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,974
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,883
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,897
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,898
; EARLIER FILING DATE: 1997-06-06

```

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; EARLIER APPLICATION NUMBER: 60/048,962
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,963
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,877
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,878
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/070,923
; EARLIER FILING DATE: 1997-12-18
; EARLIER APPLICATION NUMBER: 60/092,921
; EARLIER FILING DATE: 1998-07-15
; EARLIER APPLICATION NUMBER: 60/094,657
; EARLIER FILING DATE: 1998-07-30
; NUMBER OF SEQ ID NOS: 1227
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 416
; LENGTH: 257
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: SITE
; LOCATION: (100)
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
US-10-023-282-416
```

```

Query Match          40.3%; Score 1230; DB 14; Length 257;
Best Local Similarity 93.1%; Pred. No. 4.6e-90;
Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;
```

```

QY 160 MAALTSHLONSNNSNMNLRTRSKCKKQVFMPPSSSSSELOESRGISNFTSTHLLKEDEG 219
DB 1 MAALTSHLONSNNSNMNLRTRSKCKKQVFMPPSSSSSELOESRGISNFTSTHLLKEDEG 60
QY 220 VDDVNFRRKRPKGVITLTKGPIPKTKYKCGCKSCSGFVQSDSKRESVYCNKADAESEPEVA 279
DB 61 VDDVNFRRKRPKGVITLTKGPIPKTKYKCGCKSCSGFVQSDSKRESVYCNKADAESEPEVA 120
QY 280 QKSOLDRTVCISDAGCETLSVTSSENSLVKKKRSISGSGNPFCEKSTGIIKPKCSA 339
DB 121 QKSOLDRTVCISDAGCETLSVTSSENSLVKKKRSISGSGNPFCEKSTGIIKPKCSA 180
QY 340 KQSEHNEKEDPTFLESEIEIGTKVEVVERKEHLHTDILKRGSEMDNCSPTKDKFTGEKIF 399
DB 181 KQSEHNEKEDPTFLESEIEIGTKVEVVERKEHLHTDILKRGSEMDNCSPTKDKFT----- 235
QY 400 QEDTTPRTQIERKRTSLYF 418
DB 236 -EDTTPRTQIERKRTSLYF 253
```

```

RESULT 9
US-10-389-853-12
; Sequence 12, Application US/10389853
; Publication No. US20030180779A1
; GENERAL INFORMATION:
; APPLICANT: Lofton-Day, Cathy E.
; TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glyco
; FILE REFERENCE: 47675-36
; CURRENT APPLICATION NUMBER: US/10/389,853
; CURRENT FILING DATE: 2003-03-14
; PRIOR APPLICATION NUMBER: 60/364,689
; PRIOR FILING DATE: 2002-03-15
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 12
; LENGTH: 202
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: NON_TER
; LOCATION: (1)..(1)
```

```

; OTHER INFORMATION: functional N-terminal deletion mutant of hMBD4 (SEQ ID NO:2); aa
; OTHER INFORMATION: no. 1 corresponds to aa no. 379 of hMBD4 protein; mutant shows en
; OTHER INFORMATION: hanced deoxycytase specificity towards CpG dinucleotide sequenc
; OTHER INFORMATION: es; see Zhu et al. Nuc. Acid Res. 28:4157-4165, 2000.
US-10-389-853-12
```

```

Query Match          36.2%; Score 1106; DB 14; Length 202;
Best Local Similarity 100.0%; Pred. No. 2.8e-80;
Matches 202; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```

QY 379 GSEMDNCSPTKDPFTEKIFQEDTTPRTQIERKRTSLYSSSKYNKALSPPRKAKKW 438
DB 1 GSEMDNCSPTKDPFTEKIFQEDTTPRTQIERKRTSLYSSSKYNKALSPPRKAKKW 60
QY 439 TTPRSPNLVQETLFHDPWKLLIATIPLNRTSGMAIPVMKFLKPYSAEVAETADWRD 498
DB 61 TTPRSPNLVQETLFHDPWKLLIATIPLNRTSGMAIPVMKFLKPYSAEVAETADWRD 120
QY 499 VSELKPLGLYDLRAKTIIVFSDEYLTQWKYPIELHIGIKYGNDSYRIFCVNEWKQVHP 558
DB 121 VSELKPLGLYDLRAKTIIVFSDEYLTQWKYPIELHIGIKYGNDSYRIFCVNEWKQVHP 180
QY 559 EDHKLNKTYHDMLNENHEKLSLS 580
DB 181 EDHKLNKTYHDMLNENHEKLSLS 202
```

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RESULT 10
US-10-389-853-4
; Sequence 4, Application US/10389853
; Publication No. US20030180779A1
; GENERAL INFORMATION:
; APPLICANT: Lofton-Day, Cathy E.
; TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glycos
; FILE REFERENCE: 47675-36
; CURRENT APPLICATION NUMBER: US/10/389,853
; CURRENT FILING DATE: 2003-03-14
; PRIOR APPLICATION NUMBER: 60/364,689
; PRIOR FILING DATE: 2002-03-15
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 4
; LENGTH: 416
; TYPE: PRT
; ORGANISM: Gallus gallus
US-10-389-853-4
```

```

Query Match          28.8%; Score 880.5; DB 14; Length 416;
Best Local Similarity 78.2%; Pred. No. 9.6e-62;
Matches 161; Conservative 20; Mismatches 21; Indels 4; Gaps 2;
```

```

QY 378 RGSEMDNCS--PFRKDPFTEKIFQ--EDTTPRTQIERKRTSLYSSSKYNKALSPPRRK 433
DB 210 RDSAADGDVSWPSPDKSFYAVQAPRGTEBSAPRTQVDRRTSPYSSSKYKELASPPRRK 269
QY 434 AFKKWTPRSPFNVOETLFHDPWKLLIATIPLNRTSGMAIPVMKFLKPYSAEVAET 493
DB 270 AFKKWTPRSPFNVOETLFHDPWKLLIATIPLNRTSGMAIPVMKFLKPYSAEVAET 329
QY 494 ADMWDVSELKPLGLYDLRAKTIIVFSDEYLTQWKYPIELHIGIKYGNDSYRIFCVNEW 553
DB 330 ADMWDVSELKPLGLYDLRAKTIIVFSDEYLTQWKYPIELHIGIKYGNDSYRIFCVNEW 389
QY 554 KQVHEDHKLNKTYHDMLNENHEKLSL 579
DB 390 KQVHEDHKLNKTYHDMLNENHEKLSL 415
```

```

RESULT 11
US-10-389-853-13
; Sequence 13, Application US/10389853
; Publication No. US20030180779A1
```

```

; GENERAL INFORMATION:
; APPLICANT: Lofton-Day, Cathy E.
; TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glyco
; FILE REFERENCE: 47675-36
; CURRENT APPLICATION NUMBER: US/10/389,853
; PRIOR FILING DATE: 2003-03-14
; PRIOR APPLICATION NUMBER: 60/364,689
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: Patent version 3.1
; SEQ ID NO 13
; LENGTH: 147
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: NON TER
; LOCATION: (1)..(1)
; OTHER INFORMATION: functional N-terminal deletion mutant of hMBD4 (SEQ ID NO:2); aa
; OTHER INFORMATION: no. 1 corresponds to aa no. 434 of hMBD4 protein; mutant shows en
; OTHER INFORMATION: hanced deglycosylase specificity towards CpG dinucleotide sequenc
; OTHER INFORMATION: es; see Zhu et al. Nuc. Acid Res. 28:4157-4165, 2000.
; US-10-389-853-13

Query Match          26.7%; Score 816; DB 14; Length 147;
Best Local Similarity 100.0%; Pred. No. 3.1e-57;
Matches 147; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 444 AAKKMTPPSPFNVAOETLFHDPWKLITATITFLNPTSGMAIPVLMKFLKYPSPAEVART 493
    1 AAKKMTPPSPFNVAOETLFHDPWKLITATITFLNPTSGMAIPVLMKFLKYPSPAEVART 60

DB 494 ADMRVSELLKELGLYDLRAKTIYKFSDEYLTQWKQYPIELHIGIKYGNDSYRIFCVNEM 553
    61 ADMRVSELLKELGLYDLRAKTIYKFSDEYLTQWKQYPIELHIGIKYGNDSYRIFCVNEM 120

QY 554 KOVHPEDHKLNKYHDMWLNHEKLSLS 580
    121 KOVHPEDHKLNKYHDMWLNHEKLSLS 147

DB 121 KOVHPEDHKLNKYHDMWLNHEKLSLS 147

RESULT 12
US-10-629-951-32
; Sequence 32, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629,951
; PRIOR FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FaSTSeq for Windows Version 3.0
; SEQ ID NO 32
; LENGTH: 126
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-10-629-951-32

Query Match          22.8%; Score 697; DB 15; Length 126;
Best Local Similarity 100.0%; Pred. No. 8.6e-48;
Matches 126; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 455 DPKLLIATITFLNPTSGMAIPVLMKFLKYPSPAEVARTADMRVSELLKELGLYDLRAK 514
```

```

DB 1 DPKLLIATITFLNPTSGMAIPVLMKFLKYPSPAEVARTADMRVSELLKELGLYDLRAK 60

QY 515 TIVKPSDEYLTQWKQYPIELHIGIKYGNDSYRIFCVNEMKQVHPEDHKLNKYHDMWLNH 574
    61 TIVKPSDEYLTQWKQYPIELHIGIKYGNDSYRIFCVNEMKQVHPEDHKLNKYHDMWLNH 120

DB 575 EKLSLS 580
    121 EKLSLS 126

RESULT 13
US-10-629-951-30
; Sequence 30, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629,951
; PRIOR FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FaSTSeq for Windows Version 3.0
; SEQ ID NO 30
; LENGTH: 119
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-10-629-951-30

Query Match          20.2%; Score 616; DB 15; Length 119;
Best Local Similarity 100.0%; Pred. No. 2.5e-41;
Matches 119; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 36 KEDVAMEIERVGEDEDEQMKIRSSCNPLDQPIASQFAGTAGECRKSVPQGMERYVK 95
    1 KEDVAMEIERVGEDEDEQMKIRSSCNPLDQPIASQFAGTAGECRKSVPQGMERYVK 60

DB 96 ORLFKGTGRFPDVYRISFOGLKFRKSSILANYLHNGGTSLSKPEDPFTVLSKRGIKSR 154
    61 ORLFKGTGRFPDVYRISFOGLKFRKSSILANYLHNGGTSLSKPEDPFTVLSKRGIKSR 119

RESULT 14
US-10-629-951-37
; Sequence 37, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629,951
; PRIOR FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FaSTSeq for Windows Version 3.0
```



```
; SEQ ID NO 37
; LENGTH: 85
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-629-951-37
```

```
Query Match      14.5%; Score 443; DB 15; Length 85;
Best Local Similarity 100.0%; Pred. No. 1,1e-27;
Matches 85; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY      69  IASAFGATAGTECKSVPCGWERVVKQRLFGKTAGRFVDYFISPOGLKFRSKSLANYL 128
Db      1  IASAFGATAGTECKSVPCGWERVVKQRLFGKTAGRFVDYFISPOGLKFRSKSLANYL 60
```

```
OY      129 HKNGETSLKPEDPFTVLSKRGIKS 153
Db      61 HKNGETSLKPEDPFTVLSKRGIKS 85
```

RESULT 15

```
US-09-967-869A-7
; Sequence 7, Application US/09967869A
; Publication No. US2003008252A1
; GENERAL INFORMATION:
; APPLICANT: WOLFE, Alan P.
; APPLICANT: URNOV, Fyodor
; APPLICANT: LAI, Albert
; APPLICANT: RASCHKE, Eva
; TITLE OF INVENTION: MODULATION OF GENE EXPRESSION USING LOCALIZATION
; FILE REFERENCE: 8325-0019 / S19
; CURRENT APPLICATION NUMBER: US/09/967,869A
; CURRENT FILING DATE: 2001-09-28
; NUMBER OF SEQ ID NOS: 17
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 7
; LENGTH: 68
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: MBD4
US-09-967-869A-7
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```
Query Match      11.7%; Score 357; DB 10; Length 68;
Best Local Similarity 98.5%; Pred. No. 6,2e-21;
Matches 67; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
OY      81  ECRKSVPCGWERVVKQRLFGKTAGRFVDYFISPOGLKFRSKSLANYLHKNGETSLKPED 140
Db      1  ECRKSVPCGWERVVKQRLFGKTAGRFVDYFISPOGLKFRSKSLANYLHKNGETSLKPED 60
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OY      141 PDFTVLSK 148
Db      61 PDFTVLSK 68
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Search completed: August 22, 2005, 10:12:39
Job time : 164 secs


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Db 301 SVTSEENSLVKKERSLSGSNFCSEOKTSGIINKFCASDSEHNEKYEPTFLESEBIGT 360
Qy 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEGKIFQEDTIPRQIERRKTSLYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEGKIFQEDTIPRQIERRKTSLYFSS 420
Qy 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFDHPMKLLIATIFLNRTSGKMAIPVLMK 480
Db 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFDHPMKLLIATIFLNRTSGKMAIPVLMK 480
Qy 481 FLEKPSAEVARTADMRDVSSELKPLGLYDLRAKTIKVSDEYLTQMKYPIELHGIGKY 540
Db 481 FLEKPSAEVARTADMRDVSSELKPLGLYDLRAKTIKVSDEYLTQMKYPIELHGIGKY 540
Qy 541 GNDSYRIFCVNEMKQVHPEDHKLKYNHDMWLNENHEKSLSS 580
Db 541 GNDSYRIFCVNEMKQVHPEDHKLKYNHDMWLNENHEKSLSS 580

RESULT 2
US-09-629-222A-2
; Sequence 2, Application US/09629222A
; Patent No. 6599700
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629, 222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463, 891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053, 936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2
; LENGTH: 580
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-629-222A-2

Query Match 100.0%; Score 3055; DB 4; Length 580;
Best Local Similarity 100.0%; Pred. No. 2.2e-292;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MGTGLESLSLGDGGAAPVTSSERLVDPDPNDLRKEDVAMELERVGEDEBOMMIKSSSE 60
Db 1 MGTGLESLSLGDGGAAPVTSSERLVDPDPNDLRKEDVAMELERVGEDEBOMMIKSSSE 60
Qy 61 CNPLLOEPIASAOGFATAGTECRKSVPCGWERVVKORLFGKTAGRPDYVFISSPGLKFRS 120
Db 61 CNPLLOEPIASAOGFATAGTECRKSVPCGWERVVKORLFGKTAGRPDYVFISSPGLKFRS 120
Qy 121 KKSILANTYHKNGETSLKPEDPFTVLSKRGIKSRKYKDCSMAALTSHLONOSNNSNMNLRT 180
Db 121 KKSILANTYHKNGETSLKPEDPFTVLSKRGIKSRKYKDCSMAALTSHLONOSNNSNMNLRT 180
Qy 181 RSKCKKOVFMPSSSSSELSQESRGLSNFTSTHLLKEDBGVDVNFVRKRPKGVATLLKG 240
Db 181 RSKCKKOVFMPSSSSSELSQESRGLSNFTSTHLLKEDBGVDVNFVRKRPKGVATLLKG 240
Qy 241 IPIKTKKGRKSCSGFVQSDSKRESVCNKADASEBPVAKSQSLDRTVCI SDAGACGETL 300
Db 241 IPIKTKKGRKSCSGFVQSDSKRESVCNKADASEBPVAKSQSLDRTVCI SDAGACGETL 300
Qy 301 SVTSEENSLVKKERSLSGSNFCSEOKTSGIINKFCASDSEHNEKYEPTFLESEBIGT 360
Db 301 SVTSEENSLVKKERSLSGSNFCSEOKTSGIINKFCASDSEHNEKYEPTFLESEBIGT 360
Qy 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEGKIFQEDTIPRQIERRKTSLYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEGKIFQEDTIPRQIERRKTSLYFSS 420
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Db 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEGKIFQEDTIPRQIERRKTSLYFSS 420
Qy 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFDHPMKLLIATIFLNRTSGKMAIPVLMK 480
Db 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFDHPMKLLIATIFLNRTSGKMAIPVLMK 480
Qy 481 FLEKPSAEVARTADMRDVSSELKPLGLYDLRAKTIKVSDEYLTQMKYPIELHGIGKY 540
Db 481 FLEKPSAEVARTADMRDVSSELKPLGLYDLRAKTIKVSDEYLTQMKYPIELHGIGKY 540
Qy 541 GNDSYRIFCVNEMKQVHPEDHKLKYNHDMWLNENHEKSLSS 580
Db 541 GNDSYRIFCVNEMKQVHPEDHKLKYNHDMWLNENHEKSLSS 580

RESULT 3
US-09-657-013-56
; Sequence 56, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CpG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Tre
; FILE REFERENCE: HO-P0189US1/09905372
; CURRENT APPLICATION NUMBER: US/09/657, 013
; PRIOR FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152, 778
; PRIOR FILING DATE: 1999-09-07
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: Patentn version 3.1
; SEQ ID NO 56
; LENGTH: 580
; TYPE: PRT
; ORGANISM: Human
US-09-657-013-56

Query Match 100.0%; Score 3055; DB 4; Length 580;
Best Local Similarity 100.0%; Pred. No. 2.2e-292;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MGTGLESLSLGDGGAAPVTSSERLVDPDPNDLRKEDVAMELERVGEDEBOMMIKSSSE 60
Db 1 MGTGLESLSLGDGGAAPVTSSERLVDPDPNDLRKEDVAMELERVGEDEBOMMIKSSSE 60
Qy 61 CNPLLOEPIASAOGFATAGTECRKSVPCGWERVVKORLFGKTAGRPDYVFISSPGLKFRS 120
Db 61 CNPLLOEPIASAOGFATAGTECRKSVPCGWERVVKORLFGKTAGRPDYVFISSPGLKFRS 120
Qy 121 KKSILANTYHKNGETSLKPEDPFTVLSKRGIKSRKYKDCSMAALTSHLONOSNNSNMNLRT 180
Db 121 KKSILANTYHKNGETSLKPEDPFTVLSKRGIKSRKYKDCSMAALTSHLONOSNNSNMNLRT 180
Qy 181 RSKCKKOVFMPSSSSSELSQESRGLSNFTSTHLLKEDBGVDVNFVRKRPKGVATLLKG 240
Db 181 RSKCKKOVFMPSSSSSELSQESRGLSNFTSTHLLKEDBGVDVNFVRKRPKGVATLLKG 240
Qy 241 IPIKTKKGRKSCSGFVQSDSKRESVCNKADASEBPVAKSQSLDRTVCI SDAGACGETL 300
Db 241 IPIKTKKGRKSCSGFVQSDSKRESVCNKADASEBPVAKSQSLDRTVCI SDAGACGETL 300
Qy 301 SVTSEENSLVKKERSLSGSNFCSEOKTSGIINKFCASDSEHNEKYEPTFLESEBIGT 360
Db 301 SVTSEENSLVKKERSLSGSNFCSEOKTSGIINKFCASDSEHNEKYEPTFLESEBIGT 360
Qy 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEGKIFQEDTIPRQIERRKTSLYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEGKIFQEDTIPRQIERRKTSLYFSS 420
Qy 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFDHPMKLLIATIFLNRTSGKMAIPVLMK 480
Db 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFDHPMKLLIATIFLNRTSGKMAIPVLMK 480
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Db      421  ||||| 480
Qy      481  ||||| 540
Db      481  ||||| 540
Qy      541  ||||| 580
Db      541  ||||| 580

RESULT 4
US-09-629-222A-24
; Sequence 24, Application US/09629222A
; Patent No. 6599700
; GENERAL INFORMATION:
; APPLICANT: Bellco, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FaalSeq for Windows Version 3.0
; SEQ ID NO 24
; LENGTH: 574
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-629-222A-24

Query Match      98.6%; Score 3011; DB 4; Length 574;
Best Local Similarity 99.0%; Pred. No. 4,7e-288;
Matches 574; Conservative 0; Mismatches 0; Indels 6; Gaps 1;

Qy      1  ||||| 60
Db      1  ||||| 60
Qy      61  ||||| 120
Db      61  ||||| 120
Qy      121  ||||| 180
Db      121  ||||| 180
Qy      121  ||||| 180
Db      121  ||||| 180
Qy      181  ||||| 240
Db      181  ||||| 240
Qy      241  ||||| 300
Db      241  ||||| 300
Qy      301  ||||| 360
Db      301  ||||| 360
Qy      361  ||||| 420
Db      361  ||||| 420
Qy      421  ||||| 480
Db      421  ||||| 480
Qy      481  ||||| 540
Db      481  ||||| 540
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Qy      481  ||||| 540
Db      475  ||||| 534
Qy      541  ||||| 580
Db      541  ||||| 580

RESULT 5
US-09-629-222A-29
; Sequence 29, Application US/09629222A
; Patent No. 6599700
; GENERAL INFORMATION:
; APPLICANT: Bellco, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FaalSeq for Windows Version 3.0
; SEQ ID NO 29
; LENGTH: 384
; TYPE: PRT
; ORGANISM: Mus musculus
US-09-629-222A-29

Query Match      43.4%; Score 1326; DB 4; Length 384;
Best Local Similarity 54.3%; Pred. No. 5,3e-122;
Matches 283; Conservative 29; Mismatches 71; Indels 138; Gaps 4;

Qy      36  ||||| 95
Db      1  ||||| 57
Qy      96  ||||| 155
Db      58  ||||| 117
Qy      156  ||||| 215
Db      118  ||||| 177
Qy      216  ||||| 275
Db      178  ||||| 228
Qy      276  ||||| 335
Db      229  ||||| 228
Qy      336  ||||| 395
Db      229  ||||| 228
Qy      396  ||||| 455
Db      229  ||||| 283
Qy      456  ||||| 515
Db      284  ||||| 343
Qy      516  ||||| 555
Db      344  ||||| 384
```

RESULT 6
US-09-205-258-416
; Sequence 416, Application US/09205258
; Patent No. 6525174
; GENERAL INFORMATION:
; APPLICANT: Young et al.
; TITLE OF INVENTION: 207 Human Secreted Proteins
; FILE REFERENCE: P2007P1
; CURRENT APPLICATION NUMBER: US/09/205,258
; EARLIER FILING DATE: 1998-12-04
; EARLIER APPLICATION NUMBER: PCT/US98/11422
; EARLIER FILING DATE: 1998-06-04
; EARLIER APPLICATION NUMBER: 60/048,885
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,375
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,881
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,880
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,896
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,020
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,876
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,895
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,884
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,894
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,971
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,964
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,882
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,899
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,893
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,900
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,901
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,892
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,915
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,019
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,970
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,972
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,916
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,373
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,875
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,374
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,917
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,949
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,974
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,883

EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,897
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,898
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,962
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,963
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,877
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,878
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/070,923
; EARLIER FILING DATE: 1997-12-18
; EARLIER APPLICATION NUMBER: 60/092,921
; EARLIER FILING DATE: 1998-07-15
; EARLIER APPLICATION NUMBER: 60/094,657
; EARLIER FILING DATE: 1998-07-30
; NUMBER OF SEQ ID NOS: 1227
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 416
; LENGTH: 257
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: SITE
; LOCATION: (100)
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
US-09-205-258-416

Query Match 40.3%; Score 1230; DB 4; Length 257;
Best Local Similarity 93.1%; Pred. No. 8.2e-113;
Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;

QY 160 MAALTSHLQNSNNNNLRTSKCKKQVMPSSSELOESRGLSNFTSTHLLKDEG 219
DB 1 MAALTSHLQNSNNNNLRTSKCKKQVMPSSSELOESRGLSNFTSTHLLKDEG 60

QY 220 VDDVNFRRKVRKPKKKTITLLGIPKTKKGCRCSCSGFVQSDSKRESVCNKADSESPVA 279
DB 61 VDDVNFRRKVRKPKKKTITLLGIPKTKKGCRCSCSGFVQSDSKRESVCNKADSESPVA 120

QY 280 QKSQDLRTVCISDPAAGETLSTVSEENSLVKKKERSLSSGSNFCSEOKTSGIINKFCSA 339
DB 121 QKSQDLRTVCISDPAAGETLSTVSEENSLVKKKERSLSSGSNFCSEOKTSGIINKFCSA 180

QY 340 KDSHNEKYEEDTFLSESEIGTKVEVERKEHLHTDILKRGSEMDNNCSPTRKDTGKIF 399
DB 181 KDSHNEKYEEDTFLSESEIGTKVEVERKEHLHTDILKRGSEMDNNCSPTRKDTGKIF 235

QY 400 QEDTIPRTQIERRTKSLYF 418
DB 236 -EDTIPRTQIERRTKSLYF 253

RESULT 7
US-09-629-222A-32
; Sequence 32, Application US/09629222A
; Patent No. 6599700
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629,222A
; EARLIER FILING DATE: 2000-07-31
; EARLIER APPLICATION NUMBER: 09/463,891
; EARLIER FILING DATE: 2000-01-28
; EARLIER APPLICATION NUMBER: PCT/US98/15828
; EARLIER FILING DATE: 1998-07-28
; EARLIER APPLICATION NUMBER: 60/053,936
; EARLIER FILING DATE: 1997-07-28

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1 Sequence 1100, Application US/09205258
2 Patent No. 6525174
3
4 GENERAL INFORMATION:
5 APPLICANT: Young et al.
6 TITLE OF INVENTION: 207 Human Secreted Proteins
7 FILE REFERENCE: P2007P1
8 CURRENT APPLICATION NUMBER: US/09/205,258
9 CURRENT FILING DATE: 1998-12-04
10 EARLIER APPLICATION NUMBER: PCT/US98/11422
11 EARLIER FILING DATE: 1998-06-04
12 EARLIER APPLICATION NUMBER: 60/048, 885
13 EARLIER FILING DATE: 1997-06-06
14 EARLIER APPLICATION NUMBER: 60/049, 375
15 EARLIER FILING DATE: 1997-06-06
16 EARLIER APPLICATION NUMBER: 60/048, 881
17 EARLIER FILING DATE: 1997-06-06
18 EARLIER APPLICATION NUMBER: 60/048, 880
19 EARLIER FILING DATE: 1997-06-06
20 EARLIER APPLICATION NUMBER: 60/048, 896
21 EARLIER FILING DATE: 1997-06-06
22 EARLIER APPLICATION NUMBER: 60/049, 020
23 EARLIER FILING DATE: 1997-06-06
24 EARLIER APPLICATION NUMBER: 60/048, 876
25 EARLIER FILING DATE: 1997-06-06
26 EARLIER APPLICATION NUMBER: 60/048, 895
27 EARLIER FILING DATE: 1997-06-06
28 EARLIER APPLICATION NUMBER: 60/048, 884
29 EARLIER FILING DATE: 1997-06-06
30 EARLIER APPLICATION NUMBER: 60/048, 894
31 EARLIER FILING DATE: 1997-06-06
32 EARLIER APPLICATION NUMBER: 60/048, 971
33 EARLIER FILING DATE: 1997-06-06
34 EARLIER APPLICATION NUMBER: 60/048, 964
35 EARLIER FILING DATE: 1997-06-06
36 EARLIER APPLICATION NUMBER: 60/048, 882
37 EARLIER FILING DATE: 1997-06-06
38 EARLIER APPLICATION NUMBER: 60/048, 899
39 EARLIER FILING DATE: 1997-06-06
40 EARLIER APPLICATION NUMBER: 60/048, 893
41 EARLIER FILING DATE: 1997-06-06
42 EARLIER APPLICATION NUMBER: 60/048, 900
43 EARLIER FILING DATE: 1997-06-06
44 EARLIER APPLICATION NUMBER: 60/048, 901

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; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,892
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,915
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,019
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,970
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,972
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,916
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,373
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,875
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,374
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,917
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,949
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,974
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,883
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,897
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,898
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,962
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,963
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,877
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,878
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/070,923
; EARLIER FILING DATE: 1997-12-18
; EARLIER APPLICATION NUMBER: 60/092,921
; EARLIER FILING DATE: 1998-07-15
; EARLIER APPLICATION NUMBER: 60/094,657
; EARLIER FILING DATE: 1998-07-30
; NUMBER OF SEQ ID NOS: 1227
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1100
; LENGTH: 50
; TYPE: prt
; ORGANISM: Homo sapiens
US-09-205-258-1100

Query Match      8.3%; Score 254; DB 4; Length 50;
Best Local Similarity 98.0%; Pred. No. 1,2e-17;
Matches 49; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      110 FTSPPGLKFRSKSLANYLHKNGETSLKPEDFTVLSSKRGKISRYKDCS 159
Db      1 FSSPGLKFRSKSLANYLHKNGETSLKPEDFTVLSSKRGKISRYKDCS 50

RESULT 11
US-09-657-013-69
; Sequence 69, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of identifying Mutations in a Methyl-CPG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Tre
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; FILE REFERENCE: HO-P01893US1/09905371
; CURRENT APPLICATION NUMBER: US/09/657,013
; CURRENT FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152,778
; PRIOR FILING DATE: 1999-09-07
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 69
; LENGTH: 467
; TYPE: prt
; ORGANISM: Pro9
US-09-657-013-69

Query Match      7.2%; Score 220.5; DB 4; Length 467;
Best Local Similarity 25.7%; Pred. No. 1e-12;
Matches 98; Conservative 60; Mismatches 157; Indels 67; Gaps 15;

Qy      22 SSERLVDPDPNDLKEDVAMELER--VGDEDEOMAIKRS--SECNPFLQEPFLASAOFGA 76
Db      49 SSEH-QGEPADBEKADMSSESAEENLAVPESSASPQRORSVIRDRGMYEDP----- 99
Qy      77 TAGTECKRSVPCGWERVVKQRLFGKTAGRPDYVFISPOGLKFRSKSLANYLHKNGETSL 136
Db      100 -----TLPEGWTRLKKQKSGRSAGKFDVYLINENGAFRSKVELIYFQKVEDTSL 151
Qy      137 KPEDFTVLSSKRGKISRYKDCSMAALTSHLQNSNNNNLRTSRKCKOV--FMPPSS 194
Db      152 DPNDFTVL--TGKSPERRRQ-----KQPKKAKAPSSVSGRGKRPKSIKKVPRVK 204
Qy      195 SSELQESRGLSNFTSTHLIK----EDGEVDVNERKV-----RKPKQVTYILKQIP 242
Db      205 SEGQVAKRVIEK--SPGLLVMPYSGTKEASDATTSSQVLVILIRGGRKSE--TDPSAAP 262
Qy      243 IKTKKGCRCRKSQSFVSDSKRESVCKADABEAPVAKSQDLRTVCTISAGACGETLSV 302
Db      263 KKRGRKPSNVSLAAALAAKKAIV--KESSTKPLE-----TVLPKGRKTRRTISV 313
Qy      303 TSEE-----NSLVKK-----KERSLSSGNSFCSPQKTSGLINKFCASKDSHNKYE 349
Db      314 DVKDTIRPEPLTPIYIEKVMGQGNPAKSPESRSTGSKITGTLPKELQDHHNNHHNNH 373
Qy      350 DTFLESEEIGTKVVERKEHL 371
Db      374 HHSESKASATSPPEFSKONI 395

RESULT 12
US-09-657-013-70
; Sequence 70, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CPG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Tre
; FILE REFERENCE: HO-P01893US1/09905371
; CURRENT APPLICATION NUMBER: US/09/657,013
; CURRENT FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152,778
; PRIOR FILING DATE: 1999-09-07
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 70
; LENGTH: 467
; TYPE: prt
; ORGANISM: Pro9
US-09-657-013-70

Query Match      7.2%; Score 220.5; DB 4; Length 467;
Best Local Similarity 25.7%; Pred. No. 1e-12;
Matches 98; Conservative 60; Mismatches 157; Indels 67; Gaps 15;
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QY 22 SSERLVPDPNDLRKEDVAMELER---VGEDEEQMIRSR--SECNPLLOEPIASAOFGA 76
DB 49 SSEH-QPEBPADEGKADMSSEAEENILAVBESSASPKORSVLRDGRPMVEDP----- 99
QY 77 TAGTCRSKSVPGMGRVVKQRLFGKTAGRPDVYFISPOGLKFRSKSLANYLHNKGETSL 136
DB 100 -----TLPEGMTRKCLKQKRSKRSAGFDVYLINPGKARSVELLAIYFOKVGDSL 151
QY 137 KPEDFTVLSKRGISRYKDCSMALITSHLQNSNNMNLRTSRCKKQDV--FMPPSS 194
DB 152 DPNDPFTV--TGRGSPSRREQ-----KQPKPKAPKPSVSRGGRKRGKSIKXKPKPVK 204
QY 195 SSELQESRGLSNFTSTHLLK-----EDEGVDVNFVKY-----RKPKGVTLIKGIP 242
DB 205 SEGVOVKRIEYK-SPOKLLVKNPYSGTKEASDATTSQVLVIKRGGRKRSKSE--TDPASAP 262
QY 243 IKTKKCGKRSKSGVQSDSKRESVCNKADASEFVAOKSOLDRTVCISDAGAGETLSV 302
DB 263 KRGGRKPSNVSLAAAAAEAAKKKAI--KESIKPLLE-----TVLPKIKRKTRETTISV 313
QY 303 TSEE-----NSLVKK-----KERSLSSGSNFCSEQKTSGLINKFCSAKDSEHNEKYE 349
DB 314 DYKDTIKPEBPLTVIEKVMKGNPKAKSPBSRSTBGSPIKTLGPKKELQOHNNHHHHHH 373
QY 350 DTFLESEIIGTKVEEVERKEHL 371
DB 374 HHSESASATSPPEPETSMDNI 395

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RESULT 13

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US-09-657-013-74
; Sequence 74, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of identifying mutations in a methyl-CpG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Tre
; FILE REFERENCE: HO-P01893US1/09905371
; CURRENT APPLICATION NUMBER: US/09/657,013
; PRIOR FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152,778
; PRIOR FILING DATE: 1999-09-07
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 74
; LENGTH: 486
; TYPE: PRT
; ORGANISM: Human
US-09-657-013-74

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Query Match 7.0%; Score 213.5; DB 4; Length 486;
 Best Local Similarity 24.4%; Pred. No. 5.3e-12;
 Matches 106; Conservative 53; Mismatches 147; Indels 129; Gaps 18;

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QY 37 EDVAMELERVGEDEEQMIRKRSSECNPLLOEPIASAOFG-----ATACT 80
DB 22 KKKPLFKYKKDKKEKEGKHEPVQSAHHSABEAEAGKAEETSEGGSAAPAVEKASAP 81
QY 81 ECKRSV-----PCGWERVVKQRLFGKTAGRPDVYFISPOGLKFRSKSLANY 127
DB 82 KQRRSIIIRDGPVYDDPTLPEGWTRCLKQKRSKRSAGKTDVYLINQGAFRSKVELLIY 141
QY 128 LHNGETSLKPEDFTVLSKRGISRYKDCSMALITSHLQNSNNMNLRTSRCKKQDV 187
DB 142 FEKVGDTSIDPNDPFTV--TGRGSPSR-----REQKPPK- 175
QY 188 VFMPPSSSELQESRGL---SNFTSTHLLKDEGVDVNFVKVRKPKGVTLIKGIPRK 244
DB 176 ---PKSPKAPGTRGRGRPKSGGTTTRPKAATSEGVQVK--RVLEKSPGK--LLVQMPF- 226

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QY 245 KTKGCKRSKSGFVQSDS---KRESVCNKADASEFVAOKSOLDRTVCISDAGAGETL 300
DB 227 QTSFGAEGAGGATTSQVMVYIKRPGKRIKQADPQAIPPKR-----GKPKSVV 276
QY 301 SVTSEENSLVKKERSLSGSNFCSEQKTSGLINKFCSAKDSEHNEKEDTFLESEIIGT 360
DB 277 AAAAEE---AKKK---AVKSSIRSVOETVLPIKK-----RKTRER-----V 312
QY 361 KVEEVERKEHLHTDIL---KRGSEMDNNSPTRKDPTEBKIFQEDTTPRTQIERKTSLYF 418
DB 313 SIEVKEVVKPILVSTLAEKSGKGLTKCKSPGRK-----SKESPGR----- 354
QY 419 SSKTNKEALSPPRK 433
DB 355 ---SSASAPPKKE 365

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RESULT 14

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US-09-657-013-75
; Sequence 75, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of identifying mutations in a methyl-CpG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Tre
; FILE REFERENCE: HO-P01893US1/09905371
; CURRENT APPLICATION NUMBER: US/09/657,013
; PRIOR FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152,778
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 75
; LENGTH: 476
; TYPE: PRT
; ORGANISM: Human
US-09-657-013-75

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Query Match 6.8%; Score 209; DB 4; Length 476;
 Best Local Similarity 24.0%; Pred. No. 1.4e-11;
 Matches 104; Conservative 53; Mismatches 149; Indels 128; Gaps 17;

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QY 37 EDVAMELERVGEDEEQMIRKRSSECNPLLOEPIASAOFG-----ATACTE 81
DB 13 KDKPLKFKYKKDKKEKEGKHEPVQSAHHSABEAEAGKAEETSEGGSAARLCEASAPK 72
QY 82 CRKSV-----PCGWERVVKQRLFGKTAGRPDVYFISPOGLKFRSKSLANYL 128
DB 73 QRRSIIIRDGPVYDDPTLPEGWTRCLKQKRSKRSAGKTDVYLINQGAFRSKVELLIYF 132
QY 129 HKNGETSLKPEDFTVLSKRGISRYKDCSMALITSHLQNSNNMNLRTSRCKKQDV 188
DB 133 EKVGDTSIDPNDPFTV--TGRGSPSR-----REQKPPK- 165
QY 169 FMPPPSSSELQESRGL---SNFTSTHLLKDEGVDVNFVKVRKPKGVTLIKGIPRK 245
DB 166 ---PKSPKAPGTRGRGRPKSGGTTTRPKAATSEGVQVK--RVLEKSPGK--LLVQMPF-Q 217
QY 246 TKKGRKSCSGFVQSDS---KRESVCNKADASEFVAOKSOLDRTVCISDAGAGETLS 301
DB 218 TSPGKAEGGATTSQVMVYIKRPGKRIKQADPQAIPPKR-----GRKPGSVYA 267
QY 302 VTSSENSLVKKERSLSGSNFCSEQKTSGLINKFCSAKDSEHNEKEDTFLESEIIGT 361
DB 268 AAAAEE---AKKK---AVKSSIRSVOETVLPIKK-----RKTRER-----VS 303
QY 362 VEVVERKEHLHTDIL---KRGSEMDNNSPTRKDPTEBKIFQEDTTPRTQIERKTSLYFS 419
DB 304 IVEVKEVVKPILVSTLAEKSGKGLTKCKSPGRK-----SKESPGR----- 344

```

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QY      420 SKYKKEALSPPRK 433
      : | | | : :
Db      345 ---SSSASSPPKKE 355
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RESULT 15

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US-09-657-013-63
; Sequence 63, Application US/09657013
; Patent No. 6'709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CpG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Tre
; FILE REFERENCE: HO-P01893US1/09905371
; CURRENT APPLICATION NUMBER: US/09/657, 013
; CURRENT FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152,778
; PRIOR FILING DATE: 1999-09-07
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 63
; LENGTH: 477
; TYPE: PRT
; ORGANISM: Human
US-09-657-013-63

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Query Match	6.8%	Score 208.5	DB 4	Length 477
Best Local Similarity	23.9%	Pred. No. 1.6e-11		
Matches 104; Conservative	53;	Mismatches 149;	Indels 129;	Gaps 17;

QY	37	EDVAHELEVEVGDDEBQMTIKRSECNPLIOETIASAQFG-----	ATAGT	80
Db	13	KKPLAKFKKKVKDKKEKEKEKEPVOPSAHSHSAEPBAGCAETSESGSAPVAPEKASP	72	
QY	81	ECRKSIV-----PCGWERVVKORLPFKTAGREFDVYETISPOGLFKRSKSLIANY	127	
Db	73	KQRRSIIIRRGMYDDPPLPEGGTRTLKQKRSRSRSGKTDVYLINQKAFRSKVELIAY	132	
QY	128	LHKNGETSLKPEDPFTVLSSKGIKSRYYDCSMALLTSHLOQNSNNMNLFRSKCKCD	187	
Db	133	FEKVADTSLDPPMDPFTV-TGRGSPSR-----RQKPPKK-166		
QY	188	VEMPSSSESLQESRCL--SNFTSTHLLKEDBGVDVNFVKRKKPGKVTLLKICLPIK	244	
Db	167	----FKSPRAPGTGKRGKPKSGTTRPKPAATSEGVQV--KVLKSPKK--LIVMPF-	217	
QY	245	KTKYKCGRKSQSGFVQSDS---KRESVCNKADAESEPAQKSOQLDRTVCLISDAGAGETLU	300	
Db	218	QTSPPGAKABGCGATSTQVMVIKRPFGRKKRAEADPOAIPKR-----GRKGSVAV	267	
QY	301	SVTSESNLSVKKKESLSSGSNPFCSQKSTSGIINKRCSAKDSEHNKEYEDTPLESEIGT	360	
Db	268	AAAAABAKKAAVYESIR-----SVQETVLPIKK-----RKRET-----V	303	
QY	361	KVEVVERKEHHTDIL--KRGSEMDNNGCPTRKDFTEGKIFQEDTIPROIERRKSTSLYF	418	
Db	304	SIEVEBVAVPLIVSTLGEKSGKGLKTKCSBPGR-----SKESSPPKGR-	345	
QY	419	SSKYNKKAALSPPRK	433	
Db	346	----SSSASSPPKKE	356	

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Job time : 44 secs